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**Kasahara et al.**(10) **Patent No.:** US 9,353,379 B2  
(45) **Date of Patent:** May 31, 2016(54) **METHOD FOR CULTIVATION OF GENETICALLY-MODIFIED PLANT**(75) Inventors: **Saori Kasahara**, Tokyo (JP); **Masafumi Wasai**, Tokyo (JP); **Koichi Sugita**, Tokyo (JP); **Teruhisa Shimada**, Tokyo (JP)(73) Assignee: **NIPPON PAPER INDUSTRIES CO., LTD.**, Tokyo (JP)

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**A01H 5/10** (2006.01)  
**C12N 15/00** (2006.01)(52) **U.S. Cl.**CPC ..... **C12N 15/8261** (2013.01); **C12N 15/8234** (2013.01); **C12N 15/8238** (2013.01); **C12N 15/8251** (2013.01); **C12N 15/8257** (2013.01)(58) **Field of Classification Search**

None

See application file for complete search history.

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(57) **ABSTRACT**

The present invention provides a method for cultivation of a genetically-modified plant that can highly produce a desired protein. More specifically, the present invention provides a method for cultivation of a genetically-modified plant comprising: cultivating the genetically-modified plant in a medium,

wherein the genetically-modified plant is transformed by introducing an expression vector comprising a promoter regulating expression of RNA and a seed storage protein isolated from a plant that highly expresses RNA and the seed storage protein under a high nitrogen condition; and a polynucleotide encoding an objective protein, and

wherein the medium is adjusted so that nitrate nitrogen is 70 mg/L to 750 mg/L and/or ammonium nitrogen is 70 mg/L to 750 mg/L for a definite period in a period from 30 days before an expected flowering date to a date on or before flowering of the genetically-modified plant.

**FIG.1**

SEEDING (THE 0TH DAY)	PANICLE INITIATION STAGE (THE 50TH DAY)	HEADING, BLOOM STAGE (THE 75TH DAY)
NUTRITION GROWTH PHASE	GENERATIVE GROWTH PHASE	GRAIN FILLING PHASE
WATER 15 DAYS	CULTIVATION SOLUTION A 45 DAYS	CULTIVATION SOLUTION B 45 DAYS

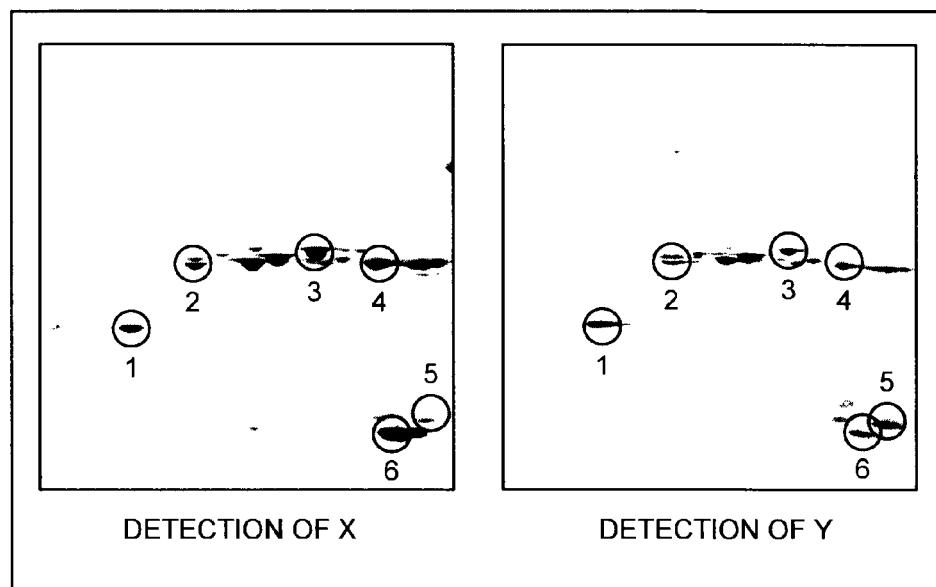
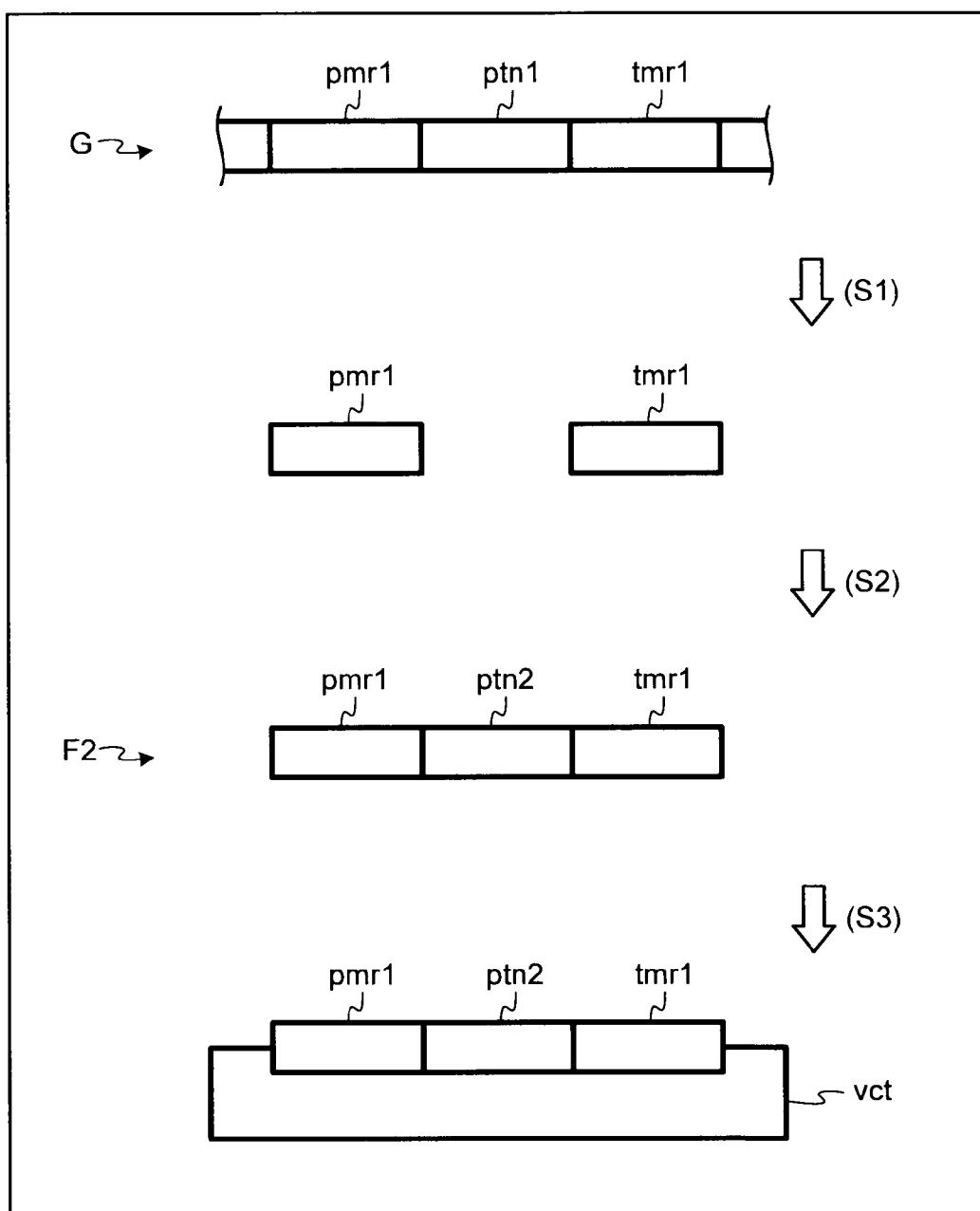
**FIG.2**

FIG.3



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**METHOD FOR CULTIVATION OF GENETICALLY-MODIFIED PLANT****TECHNICAL FIELD**

The present invention relates to a method for cultivation of a genetically-modified plant, and particularly relates to a method for cultivation of a genetically-modified plant transformed so that a desired objective protein is highly expressed, and a method for production of a seed including the desired objective protein.

**BACKGROUND ART**

Genetic modification technology has been practically applied as a breed improvement method of plants, and genetically-modified farm crops such as soybeans, maize, rapeseeds, cottons and potatoes to which functions such as herbicide resistance and harmful insect resistance had been added were developed and have been already in practical use. Further in recent years, research and development in which a useful foreign gene is introduced into a chromosome of a plant to produce a genetically-modified plant have been carried forward as a procedure to produce a functional protein or peptide for pharmaceuticals or test agents. Functional components produced by using the genetically-modified plant include not only a protein and a peptide that are a product of the introduced gene but also a product by a reaction of an introduced enzyme protein. There are many advantages in production of the functional components in a plant. The advantages include reduction of cost compared with an animal transgenic system, easy adjustment of the production scale depending on a market scale, and no possibility of contamination with a pathogen derived from an animal such as virus and prion.

As for a technology of producing efficiently functional components in a plant, for example, a method of utilizing a promoter specific for tissue in which a functional component is accumulated for genetic modification in order to control a stage, a site, and an amount of expression of a functional component has been disclosed (e.g., see Patent Document 1).

Patent Document 1: JP 2007-306941 A

**DISCLOSURE OF INVENTION****Problem to be Solved by the Invention**

However, when technology of highly accumulating a functional component in a plant body is developed, the technology of the above Patent Document 1 merely utilizes a promoter that highly expresses the functional component for genetic modification, and there is still room to improve in term of productivity of functional components.

In the light of the above circumstance, it is an object of the present invention to provide a method capable of producing a desired functional component with a higher yield by using a genetically-modified plant transformed so that the desired functional component is expressed. Any protein or peptide desired to be highly expressed in the genetically-modified plant is referred to as an "objective protein" herein in some cases.

**Means for Solving Problem**

As a result of an extensive study, the present inventors succeeded in detecting RNA and a seed storage protein that increase their expressed amounts under a predetermined high

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nitrogen-cultivation condition. Further, the present inventors succeeded in isolating a promoter that regulates expression of RNA and the seed storage protein obtained by this detection method and making a genetically-modified plant in which this promoter and a polynucleotide encoding an objective protein downstream thereof had been introduced, as well as succeeded in producing the objective protein with a high yield by cultivating such a genetically-modified plant under a predetermined high nitrogen cultivation condition. The present invention provides a method for cultivation of a following genetically-modified plant and a method for production of a seed. A term "cultivation of a plant" may be replaced by "production of a plant" herein.

[1] A method for cultivation of a genetically-modified plant comprising: cultivating the genetically-modified plant in a medium, wherein the genetically-modified plant is transformed by introducing an expression vector comprising, a promoter that regulates expression of RNA expressed in a seed that satisfies Formula (1):

$$V/W > 1.0 \quad (1)$$

wherein V is an amount of RNA contained in the seed of a predetermined plant when the plant is cultivated in a medium adjusted so that nitrate nitrogen is 70 mg/L to 750 mg/L and/or ammonium nitrogen is 70 mg/L to 750 mg/L for a definite period in a period from 30 days before an expected flowering date to a date on or before flowering, and W is an amount of RNA contained in a seed of the plant when the plant is cultivated in a medium adjusted so that nitrogen is 0 mg/L to 50 mg/L for a definite period in a period from 30 days before the expected flowering date to the date on or before flowering; and

a polynucleotide located downstream of the promoter and encoding an objective protein, and wherein the medium is adjusted so that nitrate nitrogen is 70 mg/L to 750 mg/L and/or ammonium nitrogen is 70 mg/L to 750 mg/L for a definite period in a period from 30 days before an expected flowering date to a date on or before flowering of the genetically-modified plant.

[2] A method for cultivation of a genetically-modified plant comprising: cultivating the genetically-modified plant in a medium, wherein the genetically-modified plant is transformed by introducing an expression vector comprising, a promoter that regulates expression of a seed storage protein that satisfies Formula (2):

$$X/Y > 1.0 \quad (2)$$

wherein X is an amount of the seed storage protein contained in a seed of a predetermined plant when the plant is cultivated in a medium adjusted so that a nitrate nitrogen content is 70 mg/L to 750 mg/L and/or ammonium nitrogen is 70 mg/L to 750 mg/L for a definite period in a period from 30 days before an expected flowering date to a date on or before flowering, and Y is an amount of the seed storage protein contained in a seed of the plant when the plant is cultivated in a medium adjusted so that nitrogen is 0 mg/L to 50 mg/L for a definite period in a period from 30 days before the expected flowering date to the date on or before flowering; and a polynucleotide located downstream of the promoter and encoding an objective protein, and

wherein the medium is adjusted so that nitrate nitrogen is 70 mg/L to 750 mg/L and/or ammonium nitrogen is 70 mg/L to 750 mg/L for a definite period in a period from 30 days before

an expected flowering date to a date on or before flowering of the genetically-modified plant.

[3] The method for cultivation of a genetically-modified plant according to the above [1] and [2], wherein the predetermined plant and the genetically-modified plant are the same species.

[4] The method for cultivation of a genetically-modified plant according to the above [3], wherein the predetermined plant is a poaceous plant and the genetically-modified plant is a poaceous plant.

[5] The method for cultivation of a genetically-modified plant according to the above [4], wherein the promoter is a promoter that regulates the expression of the seed storage protein selected from the group consisting of glutelin, globulin and prolamin.

[6] The method for cultivation of a genetically-modified plant according to any one of the above [1] to [5], wherein the cultivation of said genetically-modified plant is performed by water cultivation.

[7] A method for production of a seed comprising cultivating the genetically-modified plant according to any one of the above [1] to [6] and collecting the seed.

[8] The method for production of a seed according to the above [7], wherein the plant is a rice plant and the seed is a rice seed.

#### Effect of the Invention

According to the present invention, an objective protein can be produced with a high yield.

#### BRIEF DESCRIPTION OF DRAWINGS

FIG. 1 depicts a view of an outline of schedule including cultivation management;

FIG. 2 depicts a view of an image analysis result of two dimensional gel electrophoresis of the protein; and

FIG. 3 depicts a schematic view of a process of construction for an expression vector.

#### BEST MODES FOR CARRYING OUT THE INVENTION

In a method for cultivation of a genetically-modified plant of the present invention, the genetically-modified plant comprising a predetermined feature is cultivated under a predetermined condition. Embodiments of the present invention are described in detail as follows:

1. Method for making the genetically-modified plant;
2. Method for cultivation of the genetically-modified plant and method for production of its seed.

##### 1. Method for Making a Genetically-Modified Plant

The method for making a genetically-modified plant used in the present invention can be divided into following two steps as the outline, Step A and Step B. Step A is the step of detecting RNA and a seed storage protein that are highly expressed under a high nitrogen cultivation condition in a predetermined plant. Step B is the step of making a genetically-modified plant by isolating a promoter that regulates expression of RNA and the seed storage protein that are highly expressed under a high-nitrogen cultivation condition, and preparing an expression vector comprising a polynucleotide encoding an objective protein, and then introducing the expression vector into a host plant cell.

##### Step A: Detection of RNA

In the detection of RNA in Step A, a plant is cultivated under at least two different nitrogen conditions, and RNA that

is highly expressed in a certain plant species under a predetermined high nitrogen-cultivation condition is detected by using a difference of RNA contents as the indicator.

In an embodiment of method of detecting RNA in Step A, RNA that satisfies the following Formula (1) is detected.

$$V/W > 1.0 \quad (1)$$

In Formula (1), V is an amount of RNA contained in a seed of a predetermined plant when the plant is cultivated in a medium adjusted so that nitrate nitrogen is 70 mg/L to 750 mg/L and/or ammonium nitrogen is 70 mg/L to 750 mg/L for a definite period in a period from 30 days before an expected flowering date to a date on or before flowering. W is an amount of RNA contained in a seed of the plant when the plant is cultivated in a medium adjusted so that nitrogen is 0 mg/L to 50 mg/L for a definite period in a period from 30 days before the expected flowering date to the date on or before flowering.

RNA satisfying the above Formula (1) is the RNA that is highly expressed under a high nitrogen-cultivation condition. An expression regulatory region such as a promoter that facilitates expression of RNA under a high nitrogen-cultivation condition may be collected from the information of the detected RNA.

##### <Step A: Detection of Seed Storage Protein>

In the detection of a seed storage protein in Step A, a plant is cultivated under at least two different nitrogen conditions, and a seed storage protein that is highly expressed in certain plant species under a predetermined high nitrogen-cultivation condition is detected by using a difference of accumulated amounts of the seed storage protein as the indicator.

In an embodiment of method of detecting the seed storage protein in Step A, the seed storage protein that satisfies the following Formula (2) is detected.

$$X/Y > 1.0 \quad (2)$$

In Formula (2), X is an amount of the seed storage protein contained in a seed of a predetermined plant when the plant is cultivated in a medium adjusted so that a nitrate nitrogen content is 70 mg/L to 750 mg/L and/or ammonium nitrogen is 70 mg/L to 750 mg/L for a definite period in a period from 30 days before an expected flowering date to a date on or before flowering. Y is an amount of the seed storage protein contained in a seed of the plant when the plant is cultivated in a medium adjusted so that nitrogen is 0 mg/L to 50 mg/L for a definite period in a period from 30 days before the expected flowering date to the date on or before flowering.

A seed storage protein satisfying the above Formula (2) is the seed storage protein that is highly expressed under a high nitrogen-cultivation condition. An expression regulatory region such as a promoter that facilitates expression of the seed storage protein under a high nitrogen cultivation-condition may be collected from the information of the detected seed storage protein.

##### <Plant as Subject to be Tested>

The “predetermined plant” in Step A means a plant as a subject to be tested. A plant as the subject to be tested in Step A is a plant to search a promoter that facilitates expression of RNA and a seed storage protein, and may be appropriately selected in consideration of conditions such as the type of the genetically-modified plant to be made later. The plant involved in the detection of RNA and the seed storage protein is not particularly limited as long as a seed is formed. Examples may typically include: dicotyledonous plants such as tobacco, rapeseed, and soybean; and monocotyledonous plants such as grain crops including rice, maize, barley and wheat, and asparagus. Among them, the rice is a suitable plant

because ability of accumulating a protein in the seed is high and storage stability of the seed is good.

<Cultivation Condition>

The plant as the subject to be tested is cultivated under at least two different nitrogen conditions. As a preferable embodiment, cultivation is performed under First cultivation condition and Second cultivation condition. First cultivation condition is a condition in which a concentration of supplied nitrogen is high. On the other hand, Second cultivation condition is a condition in which a concentration of nitrogen is relatively lower than in First cultivation condition. Specifically, the following conditions may be included.

First cultivation condition: a plant is cultivated in a medium adjusted so that nitrate nitrogen is 50 mg/L to 750 mg/L and/or ammonium nitrogen is 50 mg/L to 750 mg/L, for a definite period in a period from 30 days before an expected flowering date to a date on or before flowering.

Second cultivation condition: a plant is cultivated in a medium adjusted so that nitrogen is 0 mg/L to 50 mg/L, for a definite period in a period from 30 days before an expected flowering date to a date on or before flowering.

The nitrogen condition in a medium opens from a time point of 30 days before an expected flowering date. The expected flowering date varies depending on a plant, and a time point of the expected flowering date in cultivated crops has been studied in every plant. For example, in case of rice plants, 30 days before the expected flowering date corresponds to a panicle formation stage.

The nitrogen condition in a medium is adjusted for a definite period in a period from between 30 days before an expected flowering date and a flowering date. It is important to expose a plant to a high nitrogen condition at least once during the period and give a definite physiological stimulation to the plant. That is, the "definite period" referred to here may be a period enough to give the definite physiological stimulation to the plant. The period of the high nitrogen condition may be appropriately adjusted depending on conditions such as a growing state and a type of a plant. Although conditions such as the growing state and the type of a plant may cause differences, it is preferable to cultivate a plant under a high nitrogen condition preferably for about one week, more preferably for about two weeks. It is also preferable to place a plant under a high nitrogen condition for a definite period immediately after passing 30 days before an expected flowering date and/or just before a flowering date. In case of rice plants, the plant may be placed under a high nitrogen-cultivation condition throughout the period from 30 days before an expected flowering date to a flowering date.

In cultivation of the plant, the nitrogen concentration of a medium is easily changed due to factors such as absorption of nutrients by the plant and an ability of a medium to retain fertilizers. In the above First and Second cultivation conditions, it is not always necessary to keep the nitrogen concentration constant in the medium, and fertilizer application may be conducted so that the nitrogen concentration falls in the above concentration range. Timing and a frequency of the fertilizer application and a concentration of the fertilizer may be appropriately adjusted as long as the nitrogen concentration in a medium is controlled as described above.

Measurement of the nitrogen concentration in a medium may be carried out, depending on the type of the medium, according to general soil analysis or fertilizer analysis methods such as the fertilizer analysis method determined by the Ministry of Agriculture, Forestry and Fisheries, and using a nitrate ion meter and an ammonia nitrogen meter.

Nitrate nitrogen refers to a nitrogen component present in a form of oxidized nitrogen such as nitrate ion. Usually,

nitrate nitrogen is present in the form of a nitrate salt obtained by binding a metal to nitrate ion in the form of  $\text{NO}_3^-$ .

Ammonium nitrogen refers to a nitrogen component present in a form of an ammonium salt among the nitrogen components.

Adjustment of a nitrogen source in a medium is conducted so that a nitrate nitrogen content in the medium is 70 mg/L to 750 mg/L and/or an ammonium nitrogen content in the medium is 70 mg/L to 750 mg/L. That is, both nitrate nitrogen and ammonium nitrogen may be used as the nitrogen sources and their concentrations may be adjusted, respectively, or either one may be used as the nitrogen source and its concentration may be adjusted. Preferably, the both are used as the nitrogen sources. A ratio of contents of the nitrate nitrogen to the ammonium nitrogen is, for example, 750:0 to 0:750, preferably 100:1 to 1:100 more preferably 30:1 to 1:30, more preferably 10:1 to 1:10, and still more preferably about 3:1 to 1:3.

A content of the nitrate nitrogen in First cultivation condition is 70 mg/L to 750 mg/L, preferably 100 mg/L to 700 mg/L and more preferably 150 mg/L to 700 mg/L. When the content of nitrate nitrogen is adjusted to 70 mg/L or less, the content of a seed storage protein reduces and it tends to be difficult to obtain an adequate indicator for comparing with that of the Second cultivation condition. On the other hand, when the content of nitrate nitrogen is adjusted to 750 mg/L or more, root rot easily occurs and causes poor growth.

A content of the ammonium nitrogen in First cultivation condition is 70 mg/L to 750 mg/L, preferably 100 mg/L to 700 mg/L and more preferably 150 mg/L to 700 mg/L. When the content of ammonium nitrogen is adjusted to 70 mg/L or less, the content of a seed storage protein reduces and it tends to be difficult to obtain an adequate indicator for comparing with that of Second cultivation condition. On the other hand, when the content of ammonium nitrogen is adjusted to 750 mg/L or more, the root rot easily occurs and causes the poor growth.

Except applying the predetermined different conditions of nitrogen as described above, fertilizer application suitable for a plant may be conducted depending on the type of a plant for cultivation. Other components contained in the medium may include phosphorous, potassium, manganese, boron, iron, calcium, copper, zinc and magnesium.

The cultivation of a plant may include a water cultivation and a soil cultivation. The soil cultivation has an advantage that considerable fluctuation of fertilizer components can be prevented because the fertilizer components are absorbed in the soil; on the flip side, the fertilizer components that can be utilized by the plant is decreased. On the other hand, the water cultivation has a feature that all of the fertilizer components in a medium can be utilized by a plant. In a selection way for plants of the present invention, a fertilizer application needs to be conducted so that a nitrogen content is adjusted under the predetermined condition, the water cultivation is more preferable in terms of easy adjustment of the fertilizer components.

<Measurement of RNA Contained in Seed and Determination of RNA>

An amount of RNA contained in a seed obtained from each plant cultivated under First and Second cultivation condition is measured. The amount of RNA contained in the seed obtained from the plant cultivated under First cultivation condition is designated as V. The amount of RNA contained in the seed obtained from the plant cultivated under Second cultivation condition is designated as W.

The above RNA amounts V and W can be obtained by utilizing various publicly known RNA detection methods. As

one embodiment, for example, RNA amount may be measured by utilizing a microarray as follows. First, RNA is extracted from a seed, and fluorescence-labeled cDNA is synthesized and then hybridized with DNA fragments on the microarray. A microarray image is imported using a scanner, and then fluorescence intensity in each spot was calculated using analysis software. The RNA amount can be obtained from the calculated spot intensity. As the seed from which RNA is extracted, a seed from 15 days to 25 days after flowering is preferable because RNA is actively synthesized.

RNA subjected to the measurement may be RNA contained in a seed. RNA for obtaining a V/W value may include, for example in case of rice plants, AK101497 (SEQ ID NO:5), AK120826, AJ002893 (SEQ ID NO:6), Os05g0329200 (SEQ ID NO:7), AK107271 (glutelin A-3) (SEQ ID NO:8), AK102194 (SEQ ID NO:9), AK120697 (SEQ ID NO:10), AK067141 (SEQ ID NO:11), AK065009 (SEQ ID NO:12), AF017360 (SEQ ID NO:13), AK071205 (SEQ ID NO:14), AK059164 (SEQ ID NO:15), AK107238 (SEQ ID NO:16), AB016505 (SEQ ID NO:17), AK099086 (SEQ ID NO:18), AY166458 (SEQ ID NO:19), AY987390 (glutelin B-2) (SEQ ID NO:20), AK107314 (SEQ ID NO:21), X15833 (SEQ ID NO:22), AY196923 (glutelin B-5) (SEQ ID NO:30), AK061894 (SEQ ID NO:23), AK107343 (glutelin 8-1) (SEQ ID NO:24), AK063995 (SEQ ID NO:25), J100041C23, AK108210 (SEQ ID NO:26), J090009107 (SEQ ID NO:27), AK065456 (SEQ ID NO:28), AK107314, AK107633 (SEQ ID NO:29), U43530 (SEQ ID NO:31), AK064310 (SEQ ID NO:32), AK064485 (SEQ ID NO:33), AK101309 (SEQ ID NO:34), AK107983 (SEQ ID NO:35), AK099918 (SEQ ID NO:36), AK100306 (SEQ ID NO:37), X83434 (SEQ ID NO:38), AK070414 (SEQ ID NO:39), AK103220 (SEQ ID NO:40), AK121856 (SEQ ID NO:41), AK062758 (SEQ ID NO:42), AK103306 (SEQ ID NO:43), AK061207 (SEQ ID NO:44), AK068266 (SEQ ID NO:45), Os06g0598500 (SEQ ID NO:46), AK119900 (SEQ ID NO:47), L19598 (SEQ ID NO:48), AK070851 (SEQ ID NO:49), Os01g0840300 (SEQ ID NO:50), AK059805 (SEQ 5 ID NO:51), AK106244 (SEQ ID NO:52), AK108127 (SEQ ID NO:53), AK108230, AK061237 (SEQ ID NO:54), AK062517 (SEQ ID NO:55), AK065259 (SEQ ID NO:56), AK065604 (SEQ ID NO:57), AK106964 (SEQ ID NO:58), AK060983 (SEQ ID NO:59), J075074G08, AK099719 (SEQ ID NO:60) (GenBank Accession No.), glutelin, globulin, and prolamin.

Based on the RNA amounts V and W, RNA that satisfies the following Formula (1):

$$\frac{V}{W} > 1.0 \quad (1)$$

is selected. Satisfying Formula (1) has a high probability that such selected RNA is RNA of which expression is increased under a predetermined high nitrogen cultivation condition. The promoter of DNA encoding such RNA has a high probability that the promoter facilitates the expression of a protein under the high nitrogen cultivation condition. The V/W value is preferably 1.25 or more, more preferably 1.5 or more and still more preferably 2.0 or more. On the other hand, when the V/W value is 1 or less, the effect of highly producing an objective protein can not be expected even if the amount of nitrogen in a medium is increased.

#### <Measurement of Content of Seed Storage Protein and Determination of Seed Storage Protein>

A content of a seed storage protein stored in the seed obtained from each plant cultivated under First and Second cultivation condition is measured. The content of the seed storage protein stored in the seed obtained from the plant cultivated under First cultivation condition is designated as X.

The content of the seed storage protein stored in the seed obtained from the plant cultivated under Second cultivation condition is designated as Y.

The above contents X and Y of the seed storage proteins may be obtained by utilizing various publicly known protein detection methods. As one embodiment, for example, a content of the protein may be measured by utilizing an electrophoresis method as follows. First, each seed storage protein is separated into a single spot on a gel by a two dimensional gel electrophoresis method. Subsequently, information on the gel is exchanged to an image file using the scanner, and then the fluorescence intensity of each spot is calculated using analysis software. A content of the protein can be obtained from the calculated spot intensity. As the seed from which a protein is extracted, a seed after 30 days, preferably 45 days after flowering is desirable because the seed storage protein is sufficiently accumulated.

The protein subjected to the measurement may be any seed storage protein. The seed storage protein for obtaining an X/Y value may include, for example in case of rice plants, glutelin, globulin and prolamin proteins, and more specifically may include glutelin B-1, glutelin B-2, glutelin B-5, glutelin A-3, globulin and 13 kDa prolamin.

Based on the contents of the seed storage protein X and Y, a seed storage protein that satisfies the following Formula (2):

$$\frac{X}{Y} > 1.0 \quad (2)$$

is selected. Satisfying Formula (2) has a high probability that the selected seed storage protein is a seed storage protein of which the expression is increased under a predetermined high nitrogen cultivation condition. The promoter of such a seed storage protein has a high probability that the promoter facilitates the expression of a protein under a high nitrogen cultivation condition. The X/Y value is preferably 1.5 or more, and more preferably 2.0 or more. On the other hand, when the X/Y value is 1 or less, the effect of highly producing an objective protein can not be expected even if the amount of nitrogen in a medium is increased.

As described above, RNA and a seed storage protein, the expression of which are increased under a predetermined high nitrogen cultivation condition, can be selected. A promoter of the selected RNA and seed storage protein facilitates expression of the protein under a high nitrogen cultivation condition. Such a promoter can be a material for a genetically-modified plant in which an objective protein is highly expressed under a predetermined high nitrogen cultivation condition by introducing the promoter into the genetically-modified plant described below.

#### <Step B: Preparation of Transformant>

In Step B, first, a promoter regulating expression of RNA and a seed storage protein that are highly expressed under a high nitrogen cultivation condition is isolated. Such a promoter may be isolated from the plant that is the subject to be tested in the above Step A and in which RNA and the seed storage protein that are highly expressed under the high nitrogen cultivation condition have been detected. The promoter may be isolated from an expression regulatory region located upstream of the nucleic acid encoding RNA and a seed storage protein.

Various RNA(s) and seed storage proteins and various nucleic acid sequences encoding them are known, shown in the known databases and the like. Also, a location and sequence of the promoter that regulates expression of RNA and a seed storage protein are publicly known in some cases. In the present invention, a promoter may be identified by utilizing publicly known sequences, or a nucleic acid sequence encoding the RNA and the seed storage protein

selected in Step A is searched and the nucleic acid sequence upstream thereof may be used as the promoter. The promoter may be identified by using the sequence commonly observed in the promoters such as TATA box, CCAAT box or a GC-rich sequence located upstream of a transcription region as a clue, or the promoter may be identified by ligating the nucleic acid encoding a known protein to a nucleic acid sequence presumed to be the promoter and measuring expression of the known protein.

<Construction of Expression Vector>

An expression vector for expressing an objective protein is constructed. The expression vector comprises at least a promoter identified as above and a nucleic acid encoding the objective protein. In addition, the others such as a terminator, a publicly known expression promoting sequence, and a marker sequence may be inserted in the expression vector.

The promoter identified as above is prepared so as to be incorporated in a vector. The promoter may be isolated from a cell of a selected plant, or may be synthesized by identifying its sequence. The vector suitable for the introduction into a plant cell is preferable. A publicly known vector or a commercially available vector may be utilized.

An objective protein may be any protein or peptide scheduled to be highly expressed in a seed. One preferable embodiment may include a so-called functional protein. The functional protein means a protein useful for human beings, e.g., antimicrobial components and enzymes. A nucleic acid encoding the objective protein may be obtained by a technique such as cloning of cDNA or genomic DNA. Also, if its DNA sequence has been previously demonstrated, the nucleic acid may be obtained by chemically synthesizing the sequence. Further, even when the DNA sequence is not demonstrated, if an amino acid sequence has been demonstrated, the DNA sequence deduced from the amino acid sequence may be synthesized chemically.

The expression vector may be made by arranging the nucleic acid encoding an objective protein downstream of the promoter and inserting this in a vector. A way to cleave or ligate a nucleic acid fragment at a predetermined position may be performed by utilizing publicly known restriction enzymes.

One embodiment for construction of a vector is described with reference to FIG. 3.

First, as the step (S1), a promoter (pmr1) contained in a chromosome G contained in the cell of the plant selected in Step A is isolated. The promoter (pmr1) together with a terminator (tmr1) constitutes a region that regulates the expression of a protein (ptn1). The promoter (pmr1) and the terminator (tmr1) are amplified from this chromosome by the technique such as PCR. PCR primers may be appropriately designed based on the sequences before and after the promoter (pmr1) and the terminator (tmr1), which sandwich each of the promoter (pmr1) and the terminator (tmr1).

Subsequently, as the step (S2), a nucleic acid fragment (F2) in which the nucleic acid (ptn2) encoding an objective protein is sandwiched between the amplified and obtained promoter (pmr1) and terminator (tmr1) is constructed.

Further, as the step (S3), the nucleic acid fragment (F2) obtained in the step (S2) is incorporated in a plasmid vector (vct). Thus, the expression vector may be constructed.

<Plant as Host>

As a host to which the vector prepared as above is introduced, a cell of a plant that produces an objective protein as the genetically-modified plant is used. The type of the plant as the host is not particularly limited as long as the above promoter is recognized and the objective protein can be expressed. A suitable plant as the host may be appropriately

selected in terms of production of an objective protein in consideration of conditions such as easiness of cultivation management, an environment of a cultivation place, a growing period, easiness of harvest, as well as nature, a size and a yield of the seed. An adverse effect on a gene expression system is easily avoided by assigning a host the same kind of plant as the plant from which the promoter facilitating expression of RNA and the seed storage protein, obtained in the above, is derived. Thus, one preferable embodiment may include a form in which the same plant species from which the promoter is derived is used as the host.

Examples of a plant as the host, that is, the plant scheduled to be cultivated as the genetically-modified plant may include: dicotyledonous plants such as tobacco, rapeseed and soybean; and monocotyledonous such as grain crops including rice plant, maize, barley and wheat, and asparagus. Among them, the rice plant is a suitable plant because an ability to accumulate a protein in a seed is high and storage stability of the seed is good.

<Introduction of Constructed Expression Vector into Plant Cell>

Subsequently, a transformed cell of a plant is made by using the constructed expression vector. A plant cell to be transformed as the host is preferably the plant that can be directly reproduced and can be cultivated in a large amount as the genetically-modified plant, in terms of simplification of steps.

As a method for introducing the constructed expression vector into the plant cell, for example, physical/chemical methods such as a microinjection method, an electroporation method, a polyethylene glycol method, a fusion method and a high speed ballistic penetration method may be used as the method for directly introducing into the plant or animal cell (I. Potrykus, Annu. Rev. Plant Physiol. Plant Mol. Biol., 42: 205, 1991). An indirect introduction method by introducing into the plant cell through virus or bacterium that infects a plant may also be used (I. Potrykus, Annu. Rev. Plant Physiol. Plant Mol. Biol., 42: 205, 1991). Viruses usable for this may include cauliflower mosaic virus, Gemini virus, tobacco mosaic virus and brom mosaic virus, and bacteria usable for this may include *Agrobacterium tumefaciens* and *Agrobacterium rhizogenes*.

<Redifferentiation of Transformant>

Subsequently, a genetically-modified tissue or a genetically-modified individual is cultured from the plant cell in which a foreign gene has been introduced by the above method. The cell having the introduced gene can be grown and redifferentiated by a standard method to culture the genetically-modified tissue or the genetically-modified individual, with appropriately performing a selection using expression of a specific character by the objective gene or a selection marker gene or disappearance of a specific character by deletion of the gene as the indicator. The seed is collected from the redifferentiated and obtained plant, and the genetically-modified body can be reproduced by utilizing the obtained seed.

Although being not altogether clear, a reason why productivity of the objective protein is enhanced by the present invention is estimated as follows. It may be considered that the genes involved in amino acid synthesis can be activated by supplying excessive nitrogen as a raw material of the amino acid synthesis during the generative growth phase before flowering. It may be considered that the genes involved in protein synthesis are activated by the effects of the amino acids accumulated in the plant body and the activated genes involved in amino acid synthesis during a grain-filling phase of the seed after the flowering phase. Further, it is considered

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that the objective protein is more efficiently produced by utilizing the promoter region of a more highly activated gene involved in protein synthesis compared with the other genes involved in protein synthesis.

## 2. Method for Cultivation of Genetically-Modified Plant and Method for Production of Seed

In a method for cultivation of genetically-modified plant of the present invention, the genetically-modified plant that can be prepared by above “1. Method for making genetically-modified plant” is cultivated under a predetermined condition. Specific embodiments are described later, and it is not always necessary to repeat above Step A: detection of RNA and seed storage protein and Step B: preparation of transformant, after once obtaining a genetically-modified plant transformed so that an objective protein is highly expressed under a high nitrogen cultivation condition by above “1. Method for making genetically-modified plant”. That is, after once obtaining the genetically-modified plant transformed so that an objective protein is highly expressed under a high nitrogen cultivation condition, the genetically-modified plant may be maintained according to an ordinary breeding method of the plant species, and the plant may be cultivated repeatedly.

### <Step C: Cultivation of Genetically-Modified Plant>

The genetically-modified plant made by above “1. Method for making genetically-modified plant” is cultivated in a medium adjusted so that nitrate nitrogen is 70 mg/L to 750 mg/L and/or ammonium nitrogen is 70 mg/L to 750 mg/L for a definite period in a period from 30 days before an expected flowering date to the flowering date.

Step C is the step of cultivating the genetically-modified plant under a predetermined high nitrogen cultivation condition. Above “1. Method for making genetically-modified plant” except only the part of “the second cultivation condition” (i.e., the case of low nitrogen condition) may be applied in the same manner to the embodiment of the cultivation condition in Step C. In other words, as an embodiment of the cultivation condition in Step C, the nitrogen condition as the “first cultivation condition” in above “1. Method for making genetically-modified plant” is applied in the same manner, and the other conditions for the cultivation: a form of a starting time of the predetermined nitrogen condition and its definite period, a form for controlling the nitrogen concentration in a medium, a form for measuring the nitrogen concentration in a medium, a form for adjusting the nitrogen source in a medium, a form for managing fertilizer application, and a form of the water cultivation or the soil cultivation may be applied in the same manner as in the embodiments described in the <cultivation condition> in above “1. Method for making genetically-modified plant”. The specific cultivation conditions other than the nitrogen condition may be appropriately adjusted depending on the type of a plant for the genetically-modified plant.

### <Method for Production of Seed>

The genetically-modified plant cultivated in the present invention may be utilized directly or by ingesting the introduced objective protein in a separated and purified form.

When the genetically-modified plant is a seed plant, the method for cultivation of the genetically-modified plant of the present invention is also the method for production of the seed. The seed plant may include poaceous plants, more preferably the rice plant because storage stability of the seed protein is high.

In case of producing the seed, when the seed is formed and reaches a predetermined maturity, the seed is collected. The collection referred to here is synonymous with harvest referred to fields of agriculture and gardening. Therefore, the

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collection includes not only separating a seed itself but also collecting a plant body including the seed from a cultivation place.

In the seed produced by the method for producing a seed plant of the present invention, an objective protein is highly expressed. The objective protein may be purified from the seed, or the seed itself may be directly utilized. When the objective protein is a functional protein that is ingestible by human, the seed may be simply ingested directly or by cooking it.

The promoter that facilitates high expression under a high nitrogen condition is found and the seed (or the cultivar) that produces a genetically-modified body in which the promoter has been incorporated is stored, thereby in the future, it is not necessary to repeat the detection of RNA and a seed storage protein and the production of the genetically-modified plant based on it every time. For example, once a promoter that regulates expression of RNA and the seed storage protein that can be highly expressed under a high nitrogen cultivation condition is identified for a certain plant, it is not necessary to perform the detection of RNA and a seed storage protein in the above Step A every time, and the genetically-modified plant may be made based on the finding.

For example, in case of isolating a promoter from rice plants, the isolated promoters may include, preferably promoters of AK101497, AK120826, AJ002893, Os05g0329200, AK107271 (glutelin A-3), AK102194, AK120697, AK067141, AK065009, AF017360, AK071205, AK059164, AK107238, AB016505, AK099086, AY166458, AY987390 (glutelin B-2), AK107314, X15833, AY196923 (glutelin B-5), AK061894, AK107343 (glutelin B-1), AK063995, J100041C23, J090009107, AK065456, AK107314, AK107633, U43530, AK064310, AK064485, AK101309, AK107983, AK099918, AK100306, X83434, AK070414, AK103220, AK121856, AK062758, AK103306, AK061207, AK068266, Os06g0598500, AK119900, L19598, AK070851, Os01g0840300, AK059805, AK106244, AK108127, AK108230, AK061237, AK062517, AK065259, AK065604, AK106964, AK060983, J075074G08, AK099719 (GenBank Accession No.), glutelin, globulin and prolamin. More preferably, the promoter may include the promoter of glutelin B-1, glutelin B-2, glutelin B-5, glutelin A-3, globulin or 13 kDa prolamin. Still more preferably, the promoter of glutelin B-1 or glutelin B-5 may be included. These promoters can facilitate expression of a structural gene present downstream thereof under a high nitrogen cultivation condition. When these promoters are used, the rice plant is suitable as a plant assigned as the host to be transformed. That is, the other embodiment of a method for cultivation of a genetically-modified plant of the present invention may include an embodiment in which a genetically-modified plant transformed is cultivated in a medium, wherein a genetically-modified plant is transformed by introducing an expression vector comprising the promoter that regulates expression of DNA encoding RNA contained in the seed satisfying V/W>1.0 and the promoter that regulates expression of the seed storage protein selected from glutelin, globulin and prolamin and the polynucleotide encoding an objective protein planned to be highly expressed, and wherein, the medium is adjusted so that nitrate nitrogen is 70 mg/L to 750 mg/L and/or ammonium nitrogen is 70 mg/L to 750 mg/L, for a definite period in a period from 30 days before an expected flowering date to the flowering.

Further, once the genetically-modified plant as mentioned above is made, the strain of the genetically-modified plant may be maintained according to the ordinary breeding

method for the plant species, and this may be cultivated according to the above step C.

### EXAMPLES

The present invention is described in more detail with reference to the following Examples, but the present invention is not limited thereto. In the following Examples, experimental manipulations in more detail for molecular biological techniques were performed according to Molecular Cloning (Sambrook et al., 1989) or instructions from the manufacturers, except the cases described particularly.

#### Example 1

##### 1. Cultivation of Plant

Rice was cultivated under First cultivation condition and Second cultivation condition as shown below. An outline of a schedule of the cultivation is shown in FIG. 1. Compositions of a cultivation solution A and a cultivation solution B used are shown in Table 1.

TABLE 1

TABLE FOR CULTIVATION  
SOLUTION COMPONENTS

COMPONENT	CULTIVATION SOLUTION A CONTENT [mg/L]	CULTIVATION SOLUTION B CONTENT [mg/L]
KNO <sub>3</sub>	38.9	621.8
NH <sub>4</sub> NO <sub>3</sub>	21.8	349.5
(NH <sub>4</sub> ) <sub>2</sub> SO <sub>4</sub>	8.3	132.7
Mg(NO <sub>3</sub> ) <sub>2</sub> •6H <sub>2</sub> O	35.7	571.4
Ca(NO <sub>3</sub> ) <sub>2</sub> •4H <sub>2</sub> O	47.0	752.6
NITRATE NITROGEN [mg/L]	33	600
AMMONIUM NITROGEN [mg/L]	17	200

##### <First Cultivation Condition for Measuring RNA Content V and Seed Storage Protein Content X>

Seeds of Nihonbare, which was one cultivar of the rice plant, were sterilized with hypochlorous acid and ethanol, subsequently spread uniformly in a petri dish in which sterilized water had been added, and cultured at 28°C. for 5 days after shielding light.

A urethane mat impregnated with water was paved in a raising seedling box in a room with artificial sunlight, and the budding seeds were seeded thereon, and cultured under a light condition: ambient temperature 28°C., humidity 50% and 11 hours, and a dark condition: ambient temperature 23°C., humidity 50% and 13 hours, for 15 days.

A cultivation bed of a submerging solution system (supplied from M Hydroponic Research Co., Ltd.) was filled with 100 L of the cultivation solution A (see Table 1), the obtained seedlings were planted one by one, and totally 77 seedlings were planted and cultured under the light condition: ambient temperature 28°C., humidity 50% and 11 hours, and the dark condition: ambient temperature 23°C., humidity 50% and 13 hours, for 45 days.

Additional fertilization was given so that the cultivation solution became the composition of the cultivation solution B, and the plants were cultured under the light condition: ambient temperature 28°C., humidity 50% and 11 hours, and the dark condition: ambient temperature 23°C., humidity 50% and 13 hours, for 45 days, and then the seeds were obtained.

Since the water cultivation was performed as above, a nitrogen concentration in the medium was controlled by controlling the nitrogen concentration in water in the cultivation bed. A nitrate nitrogen concentration was measured using 5 nitrate ion composite electrodes (supplied from DKK-TOA Corporation). An ammonium nitrogen concentration was measured using ammonia composite electrodes (supplied from DKK-TOA Corporation).

##### <Second Cultivation Condition for Measuring RNA Content W and Seed Storage Protein Content Y>

Seeds of Nihonbare were sterilized with hypochlorous acid and ethanol, subsequently spread uniformly in a petri dish in which sterilized water had been added, and cultured at 28°C. for 5 days after shielding the light.

A urethane mat impregnated with water was paved in the raising seedling box in the room with artificial sunlight, the budding seeds were seeded thereon, and cultured under the light condition: ambient temperature 28°C., humidity 50% and 11 hours, and the dark condition: ambient temperature 23°C., humidity 50% and 13 hours, for 15 days.

A cultivation pot was filled with 5 L of soil containing 50 mg/L of nitrogen, one obtained seedling was planted, and cultured under the light condition: ambient temperature 28°C., humidity 50% and 11 hours, and the dark condition: ambient temperature 23°C., humidity 50% and 13 hours, for 45 days.

The additional fertilization was given so that nitrogen in the soil was 50 mg/L, and the plant was cultured under the light condition: ambient temperature 28°C., humidity 50% and 11 hours, and the dark condition: ambient temperature 23°C., humidity 50% and 13 hours, for 45 days, and then the seeds were obtained.

##### 2. Analysis of RNA

##### <Preparation of Samples for Microarray>

The seeds obtained by cultivating under the above condition and on the 20th day after flowering were frozen in liquid 40 nitrogen, pulverized in a mortar, treated with Fruit-mate for RNA Purification (purchased from Takara Bio Inc.), and subsequently extracted with RNAiso Plus (purchased from Takara Bio Inc.). Subsequently, the sample was treated with Recombinant DNase I (RNase-free) (purchased from Takara 45 Bio Inc.), and purified using Oligotex™-dT30<Super>mRNA Purification Kit (From Total RNA) (purchased from Takara Bio Inc.), thus obtaining an RNA solution.

Total RNA was adjusted to 800 ng/one sample, and dispensed into each tube so that cyanine 3-CTP dye for labeling was 400 ng and cyanine 5-CTP dye for labeling was 400 ng. Cyanine-labeled cRNA was formed using Low RNA Fluorescent Linear Amplification Kit PLUS for two colors (purchased from Agilent Technologies Inc.), and purified using 55 RNeasy mini kit (purchased from Qiagen).

##### <Hybridization>

Eight hundred and twenty five ng of cDNA labeled with cyanine 3-CTP dye and 825 ng of cDNA labeled with cyanine 5-CTP dye were dispensed in each tube, treated with Gene 60 Expression Hybridization Kit (purchased from Agilent Technologies Inc.), and filled in rice oligo DNA microarray 4×44K RAP-DB (purchased from Agilent Technologies Inc.) to hybridize them.

##### <Image Analysis>

An image was scanned using the DNA microarray scanner (supplied by Agilent Technologies Inc.) to digitalize each spot. Further, RNA satisfying V/W>1.0 was detected using

the image analysis software of GeneSpring GX (supplied by Agilent Technologies Inc.). Values for spot intensity are shown in Table 2.

<Two Dimensional Gel Electrophoresis>

Isoelectric focusing electrophoresis for separation of proteins on the first dimension was performed using PROTEIN

TABLE 2

SPOT INTENSITY							
GENBANK ACCESSION No.	V/W	V	W	GENBANK ACCESSION No.	V/W	V	W
AK101497 (5)	1.127	91451	81143	AK107343 (24)	1.427	50738	35568
AJ002893 (6)	1.753	81314	46396	U43530 (31)	1.264	50377	39859
AJ002893 (6)	1.699	77978	45898	AK061894 (23)	1.212	50086	41334
Os05g0329200 (7)	1.177	75370	64061	AK064310 (32)	2.573	14139	5495
AK107271 (8)	1.557	75252	48343	AK064485 (33)	2.886	10076	3491
AK102194 (9)	1.751	74943	42797	AK101309 (34)	3.056	9545	3124
AK120697 (10)	1.215	71656	58957	AK107983 (35)	2.116	9180	4338
AK067141 (11)	1.005	70527	70180	AK064310 (32)	2.206	8272	3750
AK102194 (9)	1.748	69923	40007	AK099918 (36)	5.833	7159	1227
AK065009 (12)	1.116	68916	61749	AK100306 (37)	3.804	7106	1868
AF017360 (13)	1.118	68870	61584	AK099918 (36)	6.539	6960	1064
AK107271 (8)	1.736	68402	39408	X83434 (38)	2.155	5838	2709
AK120697 (10)	1.193	66986	56154	AK099918 (36)	6.427	5222	812.4
AK102194 (9)	1.715	66926	39028	AK070414 (39)	2.089	5169	2474
AK065009 (12)	1.092	65756	60201	AK103220 (40)	3.165	4961	1568
AK071205 (14)	1.014	65560	64659	AK121856 (41)	2.627	4631	1763
AK059164 (15)	1.135	65268	57482	AK121856 (41)	2.59	4432	1711
AK107238 (16)	1.413	64078	45359	AK070414 (39)	2.044	4322	2114
AF017360 (13)	1.006	64006	63630	AK062758 (42)	3.851	4153	1078
AB016505 (17)	1.238	64000	51698	AK103306 (43)	2.089	3874	1855
AK099086 (18)	1.014	62402	61556	AK061207 (44)	2.257	3780	1675
AK071205 (14)	1.088	61875	56864	AK068266 (45)	2.285	3746	1640
AY166458 (19)	1.269	61323	48330	Os06g059500 (46)	2.276	3662	1609
AY987390 (20)	1.222	60449	49479	AK119900 (47)	2.119	3565	1683
AK107314 (21)	1.034	59883	57936	L19598 (48)	2.881	3442	1195
X15833 (22)	1.477	59854	40533	AK070414 (39)	2.22	3230	1455
AK061894 (23)	1.175	59530	50670	AK121856 (41)	2.66	3043	1144
AK071205 (14)	1.046	59412	56780	AK070851 (49)	3.648	2948	808
AK107343 (24)	1.469	59250	40327	Os01g0840300 (50)	2.534	2705	1067
AK107314 (21)	1.074	58790	54715	AK059805 (51)	2.185	2704	1237
AK063995 (25)	1.066	58227	54628	AK106244 (52)	2.16	2628	1217
AK071205 (14)	1.059	57140	53974	AK108127 (53)	2.516	2569	1021
AK102194 (9)	1.744	56990	32685	AK070851 (49)	3.621	2524	697.3
J100041C23	1.507	56290	37364	AK059805 (51)	2.169	2511	1157
AK108210 (26)	1.128	55144	48904	AK108230	9.596	2398	249.9
AK102194 (9)	1.747	54892	31425	L19598 (48)	2.718	2389	879.2
J090009107 (27)	1.151	54746	47583	AK061237 (54)	3.027	2384	787.5
AK067141 (11)	1.029	53508	52003	AK062517 (55)	5.115	2240	437.9
AK107314 (21)	1.025	53428	52134	AK059805 (51)	2.162	2228	1031
AK065456 (28)	1.075	53355	49652	AK065259 (56)	2.361	2207	934.7
AK067141 (11)	1.024	53298	52053	AK065604 (57)	2.036	2123	1043
AK107314 (21)	1.04	52587	50542	L19598 (48)	2.792	2107	754.7
AK061894 (23)	1.241	52460	42255	AK106964 (58)	2.882	2093	726.3
AK061894 (23)	1.232	52324	42461	AK060983 (59)	2.228	2049	919.8
AK102194 (9)	1.908	51474	26985	J075074G08	2.106	2036	966.6
AK107633 (29)	1.006	51256	50971	AK099719 (60)	2.257	2034	901.2
AY196923 (30)	1.59	16127	10160				

## 3. Analysis of Protein

<Preparation of Sample for Two Dimensional Gel>

The seeds obtained by cultivating under the above First and Second cultivation conditions and on the 45th day after flowering were pulverized by a multi beads shocker (Yasui Kikai Corporation) after removing blastodiscs, and then homogenized in an extraction solution containing 8 M urea, 4% (w/v) SDS, 20% (w/v) glycerol and 50 mM phosphate buffer, thus obtaining a protein solution. The obtained protein solution was purified using ReadyPre 2-D Cleanup Kit (purchased from BIO-RAD LABORATORIES, Inc., hereinafter abbreviated as BIO-RAD Inc.), and ReadyStrip 7-10 Buffer (purchased from BIO-RAD Inc.) was added thereto. A total protein concentration was determined using RC DC Protein Assay (purchased from BIO-RAD Inc.).

<sup>50</sup> IEF cell (purchased from BIO-RAD Inc.) and 7 cm ReadyStrip IPG Strip 3-7NL (purchased from BIO-RAD Inc.). Electrophoresis for the separation of the proteins on the second dimension was performed by equilibrating IPG Strip with

<sup>55</sup> equilibration buffer I and equilibration buffer II (purchased from BIO-RAD Inc.) and using PROTEIN cell (purchased from BIO-RAD Inc.) and 10-20% resolving Ready Gel Pre-cast Gel (purchased from BIO-RAD Inc.). The samples were

<sup>60</sup> electrophoresed with molecular weight markers and isoelectric point calibration markers in order to calculate the molecular weights and the isoelectric points of protein spots upon image analysis. The gel immediately after performing the two dimensional SDS-PAGE was immersed in a fixation solution containing 40% ethanol and 10% acetic acid for 2 hours, and treated with Flamingo Gelstain (purchased from BIO-RAD Inc.).

## &lt;Image Analysis of Two Dimensional Gel&gt;

The treated gel was digitalized using Pharus FX Molecular Imager (purchased from BIO-RAD Inc.). Spot positions of 6 types of the seed storage proteins were identified and the spot intensity was measured using Quantity One and PDQuest (purchased from BIO-RAD Inc.). Six types of the seed storage proteins were globulin, glutelin, glutelin B-5, glutelin B-1, 10 kDa prolamin and 13 kDa prolamin. An image analysis view of the two dimensional gel and the values of the spot intensity are shown in FIG. 2 and Table 3, respectively.

TABLE 3

PROMOTER	SPOT INTENSITY					
	SPOT No.					
	1	2	3	4	5	6
GLOBULIN						
SPOT X	8338	4530	7902	9159	1521	16784
INTENSITY Y	7624	2369	3292	4220	1730	4587
X/Y	1.09	1.91	2.40	2.17	0.88	3.66
PROLAMIN						
10 KDa B-1						
13 KDa PROLAMIN						

## 4. Preparation of Genetically-Modified Plant

## &lt;Preparation of Expression Vector&gt;

As a result of above “2. Analysis of RNA” and “3. Analysis of protein”, a nucleic acid sequence encoding glutelin B-1 having V/W=1.47 and X/Y=2.17 (GluB-1 gene: Accession No. X54314, AK107343), and a promoter sequence thereof (GluB-1 Promoter region: Accession No. AY427569) were obtained from NCBI Home Page, Nucleotide Database. Nucleic acid sequences neighboring to the nucleic acid sequence encoding glutelin B-1 were searched by Rice Annotation Project Database, and a nucleic acid sequence of 1.0 kb downstream of the nucleic acid encoding glutelin B-1 was obtained as a terminator sequence.

PCR primers were designed based on the obtained nucleic acid sequence information. A fragment (fragment A) comprising the promoter sequence of glutelin B-1, in which an Sse8387I site had been added to its N terminus, and a signal peptide sequence of glutelin B-1 was obtained by PCR (Primer1: CCTGCAGGACAGATTCTTGCTACCAACA (SEQ ID NO:1), Primer2: CAGGAGTGGAGTATC-GAGGTAAAAGAA (SEQ ID NO:2)). A fragment (fragment B) comprising the terminator sequence on which SacI site had been added to its N terminus and EcoRI site had been added to its C terminus was also obtained by PCR (Primer3: GAGCTCTGTAATTGAGAACTAGTATC (SEQ ID NO:3), Primer4: GAATTCTCTTAACCTTACCTATGAT (SEQ ID NO:4)).

The obtained fragments A and B were introduced into *E. coli* using Zero Blunt TOPOPCR cloning kit (purchased from Invitrogen Corporation), and amplified. The fragment A, and a fragment C linking 7crp (pollen disease alleviation peptide) and a KDEL sequence (JP 2004-321079-A) were treated with NcoI (purchased from Takara Bio Inc.), and subsequently ligated using DNA Ligation Kit (purchased from Takara Bio Inc.). The fragment ligating the fragment A and the fragment C, and the fragment B were treated with SacI (purchased from Takara Bio Inc.), and ligated using DNA Ligation Kit (purchased from Takara Bio Inc.). The resulting ligated fragment (A, B, C) was treated with Sse8387I (purchased from Takara Bio Inc.) and EcoRI (purchased from Takara Bio Inc.), and subsequently ligated between restriction enzyme sites of EcoRI-Sse8387I of a plasmid pTL7 (H. Ebinuma et al.,

Molecular Methods of Plant Analysis, 22: 95, 2002) using DNA Ligation Kit (purchased from Takara Bio Inc.) to construct an expression vector. The above SEQ ID NOS correspond to those in Sequence Listing.

<Introduction into *Agrobacterium*>

*Agrobacterium tumefaciens* (*A. tumefaciens*) EHA 105 strain was inoculated in 10 mL of YEB liquid medium (5 g/L of beef extract, 1 g/L of yeast extract, 5 g/L of peptone, 5 g/L of sucrose, 2 mM MgSO<sub>4</sub>, pH 7.2 at 22° C.) (hereinafter pH is the pH at 22° C. unless otherwise specified), and cultured at

28° C. until OD<sub>630</sub> reached the range of 0.4 to 0.6. After the culture, the cultured medium was centrifuged at 6900×g at 4° C. for 10 minutes to collect microbial cells. The collected microbial cells were suspended in 20 mL of 10 mM HEPES (pH 8.0), and the microbial cells were collected again by centrifuging at 6900×g at 4° C. for 10 minutes. These microbial cells were suspended in 200 μL of the YEB liquid medium to use as a bacterial solution for introducing the plasmid. The expression vector and 50 μL of the above bacterial solution for introducing the plasmid were mixed in a 0.5 mL tube, and the expression vector was introduced into *A. tumefaciens* EHA 105 strain using an electroporation method (Gene Pulsar II System (purchased from BIO-RAD Inc.). The microbial cells after the expression vector had been introduced were added to 200 μL of the YEB liquid medium and cultured at 25° C. for one hour with shaking. Subsequently, the microbial cells were seeded on YEB agar medium (1.5 w/v % agar, the other composition was the same as above) containing 50 mg/L of kanamycin, and cultured at 28° C. for 2 days. Then, a growing bacterial colony was transferred into the YEB liquid medium and further cultured. The plasmid was extracted from the growing microbial cells by an alkali method, and it was confirmed that the expression vector was introduced into these microbial cells.

<Transformation of Rice Plant with *Agrobacterium* EHA 105 Strain>

Completely matured seeds of the rice plant cultivar, “Nihonbare” were sterilized according to the method in Experimental Protocol for Model Plant (pages 93 to 98) in Cell Engineering Supplementary Volume, Plant Cell Engineering Series 4, and subsequently placed on N6C12 medium (N6 inorganic salts and vitamins (Chu C. C., 1978, Proc. Symp. Plant Tissue Culture, Science Press Peking, pp. 43-50), 30 g/L of sucrose, 2.8 g/L of proline, 0.3 g/L of casamino acids, 2 mg/L of 2,4-D, 4 g/L of Gel-Lyte, pH=5.8), cultured in a light place at 28° C. with being sealed with a surgical tape to sprout, and used as a material to be infected with *Agrobacterium* EHA 105. *Agrobacterium* EHA 105 introduced by the expression vector cultured on the YEB agar medium (15 g/L of Bacto agar, the other composition is the same as above) was transferred to the YEB liquid medium, cultured at 25° C. at 180 rpm overnight, subsequently microbial cells were collected by centrifuging at 3000 rpm for 20

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minutes, and suspended in N6 liquid medium (N6 inorganic salts and vitamins, 30 g/L of sucrose, 2 mg/L of 2,4-D, pH=5.8) containing 10 mg/L of acetosyringone so that OD<sub>630</sub> was 0.15 to use as a suspension of *Agrobacterium* for infection. The prepared budding seeds were placed in a 50 mL tube, and the suspension of *Agrobacterium* for infection is poured on it, and the budding seeds were immersed for 1.5 minutes in the suspension of *Agrobacterium* for infection. After the immersion, the suspension of *Agrobacterium* for infection was discarded, the budding seeds were placed on a sterilized filter, and extra water was removed. These seeds were placed on the coexistence culture medium of N6C12 medium (N6 inorganic salts and vitamins, 30 g/L of sucrose, 2.8 g/L of proline, 0.3 g/L of casamino acids, 2 mg/L of 2,4-D, 4 g/L of Gel-Lyte, pH=5.2), and cultured in a dark place at 28° C. for 3 days with being sealed with the surgical tape, and then transferred to N6C12CH25 medium (N6 inorganic salts and vitamins, 30 g/L of sucrose, 2.8 g/L of proline, 0.3 g/L of casamino acids, 2 mg/L of 2,4-D, 500 mg/L of carbenicillin, 25 mg/L of hygromycin, 4 g/L of Gel-Lyte), and cultured.

## &lt;Redifferentiation of Transformant&gt;

One week after the start of the culture in the above N6C12H25 medium, a budding sprout was removed from blastodisc tissue, and the remaining blastodisc tissue was cultured in the N6C14-CH25 medium (N6 inorganic salts and vitamins, 30 g/L of sucrose, 2.8 g/L of proline, 0.3 g/L of casamino acids, 4 mg/L of 2,4-D, 500 mg/L of carbenicillin, 25 mg/L of hygromycin, 4 g/L of Gel-Lyte) for one week, and further transferred to and cultured in MSRC medium (MS inorganic salts and vitamins (Murashige, T. and Skoog, F., 1962, Physiol. Plant., 15, 473), 30 g/L of sucrose, 30 g/L of sorbitol, 2 g/L of casamino acids, 500 mg/L of carbenicillin, 4 g/L of Gel-Lyte), thus redifferentiating a sprout or a young plant body.

## 5. Cultivation of Genetically-Modified Plant

The rice plant obtained by above “3. Preparation of genetically-modified plant” was cultivated as follows. The compositions of a cultivation solution C and a cultivation solution D used for the cultivation are shown in Table 4.

First, the sprout or the young plant body redifferentiated from the blastodisc tissue obtained by above “3. Preparation of genetically-modified plant” was transferred to a taking root medium, and grown until a young seedling having a height of about 20 cm was obtained.

The cultivation bed of the submerging solution system (supplied from M Hydroponic Research Co., Ltd.) was filled with 100 L of the cultivation solution C (see Table 4), the obtained seedlings were planted one by one, totally 77 seedlings were planted and cultured under the light condition: ambient temperature 28° C., humidity 50% and 11 hours, and the dark condition: ambient temperature 23° C., humidity 50% and 13 hours, for 45 days.

Additional fertilization was given so that the cultivation solution became the composition of the cultivation solution D (see Table 4), and the plants were cultured under the light condition: ambient temperature 28° C., humidity 50% and 11 hours, and the dark condition: ambient temperature 23° C., humidity 50% and 13 hours, for 45 days.

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TABLE 4

COMPONENT	TABLE FOR CULTIVATION SOLUTION COMPONENTS	
	CULTIVATION SOLUTION C	CULTIVATION SOLUTION D
	CONTENT [mg/L]	
KNO <sub>3</sub>	38.9	174.1
NH <sub>4</sub> NO <sub>3</sub>	21.8	97.9
(NH <sub>4</sub> ) <sub>2</sub> SO <sub>4</sub>	8.3	37.1
Mg(NO <sub>3</sub> ) <sub>2</sub> •6H <sub>2</sub> O	35.7	160.0
Ca(NO <sub>3</sub> ) <sub>2</sub> •4H <sub>2</sub> O	47.0	210.7
NITRATE NITROGEN [mg/L]	33	150
AMMONIUM NITROGEN [mg/L]	17	50

## 6. Measurement of Protein Content

Seeds were collected from the plant obtained in above “4. Cultivation of genetically-modified plant”, and a protein content in the seed was measured as follows.

<Total Protein Content in Seed>

The protein content in the seed from which the blastodisc had been removed was measured using a near-infrared ray protein analysis apparatus NIRFLEX N-500 (supplied from BUCHI), and the total protein content was calculated from a seed weight.

<Amount of Functional Protein (Pollen Disease Alleviation Peptide: 7crp)>

The protein in the seed and protein markers with known concentrations were electrophoresed using SDS-PAGE kit (purchased from BIO-RAD Inc.), and then the gel was immersed in the fixation solution containing 40% ethanol and 10% acetic acid for two hours, and treated with Flamingo Gelstain (purchased from BIO-RAD Inc.). The treated gel was digitalized using Pharos FX Molecular Imager (purchased from BIO-RAD Inc.). A band position of 7crp was identified using Quantity One (purchased from BIO-RAD Inc.), and the weight of the pollen disease alleviation peptide (7crp) was calculated by comparing with the band intensity of the marker with the known concentration. In this way, the amount of the total protein in the seed collected from the genetically-modified plant and the amount of the functional protein introduced by the transformation were obtained.

## Example 2

A genetically-modified plant was cultivated and a protein content was measured in the same manner as in Example 1, except that an ammonium nitrogen content of the cultivation solution D was changed to 150 mg/L.

## Example 3

A genetically-modified plant was cultivated and a protein content was measured in the same manner as in Example 1, except that a nitrate nitrogen content of the cultivation solution D was changed to 50 mg/L and an ammonium nitrogen content of the cultivation solution D was changed to 150 mg/L.

## Example 4

A genetically-modified plant was cultivated and a protein content was measured in the same manner as in Example 1, except that a nitrate nitrogen content of the cultivation solu-

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tion D was changed to 600 mg/L and an ammonium nitrogen content of the cultivation solution D was changed to 200 mg/L.

## Example 5

A genetically-modified plant was cultivated and a protein content was measured in the same manner as in Example 1, except that a nitrate nitrogen content of the cultivation solution D was changed to 70 mg/L and an ammonium nitrogen content of the cultivation solution D was changed to 70 mg/L.

## Example 6

A genetically-modified plant was cultivated and a protein content was measured in the same manner as in Example 1, except that a nitrate nitrogen content of the cultivation solution D was changed to 750 mg/L and an ammonium nitrogen content of the cultivation solution D was changed to 750 mg/L.

## Example 7

A genetically-modified plant was cultivated and a protein content was measured in the same manner as in Example 1, except that the promoter of glutelin B-5 was used as the promoter for expression of the objective protein.

## Example 8

A genetically-modified plant was cultivated and a protein content was measured in the same manner as in Example 1, except that the cultivation condition of Y was the water cultivation.

## Example 9

A genetically-modified plant was cultivated and a protein content was measured in the same manner as in Example 1, except that an ammonium nitrogen content of the cultivation solution B was changed to 20 mg/L.

## Example 10

A genetically-modified plant was cultivated and a protein content was measured in the same manner as in Example 1, except that a nitrate nitrogen content of the cultivation solution B was changed to 20 mg/L.

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## Comparative Example 1

A genetically-modified plant was cultivated and a protein content was measured in the same manner as in Example 1, except that a nitrate nitrogen content of the cultivation solution D was changed to 20 mg/L and an ammonium nitrogen content of the cultivation solution D was changed to 20 mg/L.

## Comparative Example 2

<sup>10</sup> A genetically-modified plant was cultivated and a protein content was measured in the same manner as in Example 1, except that a nitrate nitrogen content of the cultivation solution D was changed to 800 mg/L and an ammonium nitrogen content of the cultivation solution D was changed to 800 mg/L.

## Comparative Example 3

<sup>20</sup> A genetically-modified plant was cultivated and a protein content was measured in the same manner as in Example 1, except that the promoter of 10 kDa prolamin was used as the promoter for expression of the objective protein.

## Comparative Example 4

<sup>25</sup> A genetically-modified plant was cultivated and a protein content was measured in the same manner as in Example 1, except that a nitrate nitrogen content of the cultivation solution B was changed to 20 mg/L and an ammonium nitrogen content of the cultivation solution B was changed to 20 mg/L.

## Comparative Example 5

<sup>30</sup> A genetically-modified plant was cultivated and a protein content was measured in the same manner as in Example 1, except that a nitrate nitrogen content of the cultivation solution B was changed to 800 mg/L and an ammonium nitrogen content of the cultivation solution B was changed to 800 mg/L.

## Comparative Example 6

<sup>35</sup> A genetically-modified plant was cultivated and a protein content was measured in the same manner as in Comparative Example 3, except that a nitrate nitrogen content of the cultivation solution D was changed to 20 mg/L and an ammonium nitrogen content of the cultivation solution D was changed to 20 mg/L.

Test results of above Examples 1 to 10 and Comparative Examples 1 to 6 are shown in following Table 5.

TABLE 5

	TEST RESULTS										CULTIVATION OF GENETICALLY-						
	SELECTION OF PROMOTER FOR SEED STORAGE PROTEIN GENE					MODIFIED PLANT					PROTEIN AMOUNT						
	X		Y		NITROGEN CONTENT [mg/L]	NITROGEN CONTENT [mg/L]		NITROGEN BEFORE FLOWERING [mg/L]		NITROGEN BEFORE FLOWERING [mg/L]		PROMOTER (AK107343)	NITRATE AMMONIUM NITROGEN CULTIVATION METHOD		WATER CULTIVATION	2.09 11.2	
	NITROGEN CONTENT BEFORE FLOWERING [mg/L]	NITRATE AMMONIUM NITROGEN CULTIVATION METHOD	TOTAL NITROGEN CULTIVATION METHOD	CULTIVATION METHOD		SOIL CULTIVATION	SOIL CULTIVATION	GLUTELIN B-1 (AK107343)	GLUTELIN B-1 (AK107343)	GLUTELIN B-1 (AK107343)	GLUTELIN B-1 (AK107343)		V/W X/Y NITROGEN CULTIVATION METHOD				
EXAMPLE 1	600	200	50	SOIL CULTIVATION	50	GLUTELIN B-1 (AK107343)	1.47	2.17	1.50	50	WATER CULTIVATION	2.09	11.2				
EXAMPLE 2	600	200	50	SOIL CULTIVATION	50	GLUTELIN B-1 (AK107343)	1.47	2.17	150	150	WATER CULTIVATION	2.19	11.7				
EXAMPLE 3	600	200	50	SOIL CULTIVATION	50	GLUTELIN B-1 (AK107344)	1.47	2.17	50	150	WATER CULTIVATION	1.93	10.3				
EXAMPLE 4	600	200	50	SOIL CULTIVATION	50	GLUTELIN B-1 (AK107344)	1.47	2.17	600	200	WATER CULTIVATION	2.66	14.3				
EXAMPLE 5	600	200	50	SOIL CULTIVATION	50	GLUTELIN B-1 (AK107345)	1.47	2.17	70	70	WATER CULTIVATION	1.95	10.4				
EXAMPLE 6	600	200	50	SOIL CULTIVATION	50	GLUTELIN B-1 (AK107346)	1.47	2.17	750	750	WATER CULTIVATION	2.59	13.9				
EXAMPLE 7	600	200	50	SOIL CULTIVATION	50	GLUTELIN B-1 (AK107347)	1.59	2.40	150	50	WATER CULTIVATION	2.12	12.6				
EXAMPLE 8	600	200	50	SOIL CULTIVATION	50	GLUTELIN B-1 (AY196923)	—	—	—	—	—	—	—				
EXAMPLE 9	600	20	50	SOIL CULTIVATION	50	GLUTELIN B-1 (AK107347)	—	—	—	—	—	—	—				
EXAMPLE 10	20	200	50	SOIL CULTIVATION	50	GLUTELIN B-1 (AK107347)	—	—	—	—	—	—	—				
COMPARATIVE EXAMPLE 1	600	200	50	SOIL CULTIVATION	50	GLUTELIN B-1 (AK107347)	1.47	2.17	20	20	WATER CULTIVATION	1.14	6.1				
COMPARATIVE EXAMPLE 2	600	200	50	SOIL CULTIVATION	50	GLUTELIN B-1 (AK107347)	1.47	2.17	800	800	WATER CULTIVATION	*1	—				
COMPARATIVE EXAMPLE 3	600	200	50	SOIL CULTIVATION	50	GLUTELIN B-1 (AK107347)	0.80	0.88	150	50	WATER CULTIVATION	2.01	4.4				
COMPARATIVE EXAMPLE 4	20	20	50	SOIL CULTIVATION	50	PROLAMIN GLUTELIN B-1 (AK107347)	—	0.98	—	—	—	—	—				
COMPARATIVE EXAMPLE 5	800	800	50	SOIL CULTIVATION	50	PROLAMIN GLUTELIN B-1 (AK107347)	—	*2	—	—	—	—	—				
COMPARATIVE EXAMPLE 6	600	200	50	SOIL CULTIVATION	50	PROLAMIN GLUTELIN B-1 (10 KDa)	0.80	0.88	20	20	WATER CULTIVATION	1.12	3.0				

\*1,2: NO SEED COULD BE COLLECTED DUE TO THE ROOT ROT.

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## INDUSTRIAL APPLICABILITY

The present invention is useful in fields such as biomass production, functional food production, biotechnology and breed improvement of the plants.

## REFERENCE SIGNS LIST

G: chromosome  
pmr 1: promoter

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ptn1, ptn2: a structural gene region encoding a protein  
tmr1: terminator  
vct: plasmid vector

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## SEQUENCE LISTING FREE TEXT

SEQ ID NO. 1: Primer 1  
SEQ ID NO. 2: Primer 2  
SEQ ID NO. 3: Primer 3  
SEQ ID NO. 4: Primer 4

## SEQUENCE LISTING

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120

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aatccgagca	agaccaacaa	tttgcgttcaa	aaagccaaag	ccataaaattt	agagatgaac	480
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atgggtgctt	caatgtatgg	gatgcacca	tttgcgttcat	atatgtact	gatataataca	600
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tttgcgttgc	gagggttgc	caaaatatcg	acaacccaaa	cctcgacat	acatacaacc	1020
ccagagcagg	aaggatcaca	tatctaaatg	gccaatgtt	ccccattttt	aatcttgc	1080
agatgatgtc	cgttaaagta	aatttataatc	agaacgcact	ccttcacact	ttttggaaaca	1140
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acaatggaaa	gacagtgttc	gatggagac	tccgtcggt	gcagttctt	attataaccac	1260
aacaccatgt	agtcatataaa	aaggcacaaa	gggaggat	ctcatatatt	gcattgaaaa	1320
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ctgacgtgt	tgttagcaat	gcatatcgta	tctcaagaga	agaagctt	aggctcaac	1440
acaacagggg	agatgatgttca	cttgcgttgc	tttgcgttgc	tttgcgttgc	tttgcgttgc	1500
acatatctgt	gagtgcataa	ccaagaaatg	tcaatcatag	gttggat	ctaaggatgg	1560
tttcaataa	aatcataagc	aataaaagag	tgcgttgc	tgcgttgc	taatgttgc	1620
tgggttggaa	aaataataata	aatttgcgtt	ccttt			1655

&lt;210&gt; SEQ ID NO 9

&lt;211&gt; LENGTH: 1745

&lt;212&gt; TYPE: DNA

&lt;213&gt; ORGANISM: Oryza sativa

&lt;220&gt; FEATURE:

&lt;221&gt; NAME/KEY: misc\_feature

<223> OTHER INFORMATION: LOCUS: AK102194 1745 bp mRNA linear PLN  
04-DEC-2008; DEFINITION: Oryza sativa Japonica Group cDNA  
clone:J033087C05, full insert sequence.; ACCESSION: AK102194

&lt;400&gt; SEQUENCE: 9

caagccgaga	caacgcagag	aaaggcgct	tcgtactcgc	ctctctccgc	gcctccgc	60
ttttcctcct	cctctccctt	ctctccctt	tccggccgc	tcgcagatc	aacccaaatcc	120

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ggcgccatga gggagtgcac	ctcgatccac atcggccagg	ccggtatcca ggtcgaaac	180
gctgtctggg agctctactg	cctcgagcat ggcatccagg	ctgatggta gatgccagt	240
gacaggactg ttggtgagg	tgtatgtgt ttcaacacct	tcttcagtga gactggtgc	300
ggaaagcatg ttccccgtgc	tgtatgttt gatcttgagc	ctactgtgtat tgatgagggt	360
aggactggtt gctaccgcca	gctcttccac cctgagcagc	tcatcaatgg caaggaggat	420
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gtggactatg gcaagaagtc	caagcttggg ttcaactgtgt	acccatcccc tcaggctc	660
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gatgttgctg ttctgcttga	taatgaggcc atctatgaca	tctgcccggc ctcccttgc	780
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cagtgtggag ttatcctgtt	gcccgtgtcg tgggtcttcg	agacatttc ttgtgggtat	1620
ctgatgtttt gggcttaatg	ggcttacaac ctcgttgc	gtaacctgtg tgccctgtt	1680
tatgtgagac cgcttcgtca	cttataatat ggcgtttgt	tcaattttat gtttgc	1740
tgtgt			1745

<210> SEQ ID NO 10  
 <211> LENGTH: 752  
 <212> TYPE: DNA  
 <213> ORGANISM: Oryza sativa  
 <220> FEATURE:  
 <221> NAME/KEY: misc\_feature  
 <223> OTHER INFORMATION: LOCUS: AK120697 752 bp mRNA linear PLN  
 04-DEC-2008; DEFINITION: Oryza sativa Japonica Group cDNA  
 clone:J013170I05, full insert sequence.; ACCESSION: AK120697

&lt;400&gt; SEQUENCE: 10

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cggcgaccc ctccttcgtat	ccatatcttc cggtcgagtt	cttgggtcgat ctctccctc	120
ctccacccatcc tcctcacagg	ttcttgggtgt agcttgcac	tttcaccacg aaagtttcat	180
gtctgatctc gacattcaga	tcccaactgc ctgcgttgc	tttgcgttgc ccaatgcgtt	240
agactctgtt gcggtgcag	gatcaaagga ctacgttcat	gtacgcaccc agcagcgtaa	300

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tggccgtaag agcctgacca ctgtccaggg	attgaagaag gaattcagct acaaacaagat	360
cctcaaagat ctcagaaggaaag agtttgctg	caatggact gttgtccagg acccagagct	420
tggccaggtc attcaacttc	agggtgatca gaggaaaaac gtatcaaatt ttcttgcata	480
ggccggcatt gtgaagaagg aacacatcaa	gattcatggt ttctgagcaa ctgccaaaac	540
cattgcaag actatagttt	ggggtggagt atacttggtt gtgtacatgc ctgcgtttc	600
cattgtacac acaaaaaccta	gccacctt gactctttag tttatgtctt ttaccctgt	660
gttgaagttt	gtaagaggca ccatcaactat agatgtatggc ttgtgtccct ctttcatcaa	720
gattgaataa tatatgctac	tttgagagcg ct	752

<210> SEQ ID NO 11  
<211> LENGTH: 1130  
<212> TYPE: DNA  
<213> ORGANISM: Oryza sativa  
<220> FEATURE:  
<221> NAME/KEY: misc\_feature  
<223> OTHER INFORMATION: LOCUS: AK067141 1130 bp mRNA linear PLN  
04-DEC-2008; DEFINITION: Oryza sativa Japonica Group cDNA  
clone:J013095J21, full insert sequence.; ACCESSION: AK067141

<400> SEQUENCE: 11

gatccatcca tccatgtcgt	gttgttagct tcgtttctt cttgtttcat caatccaagg	60
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ttcagtggtt	agcgttagctt gattagttgg tttaatttagt tggagagtaa agaagatgtt	180
gggagtgttc	agcggcgcacg tggggaggt gcccggcggag ctggggcccg ccggcagccg	240
gacgcgtcg	ccgaagacgc gggcgctgga gctggcgtac egcttectcg gcggcgccga	300
gccccccgtg	tccgtgcage tcggcgcacct cggccaccc tcgtactccc atgccaacca	360
agcccttc	cgtccaaaggcat catttgcagc aaaagatgac atcttctgcc tttcgaggg	420
agtcctggac	aatcttggca acctgaacca gcagttggt ctgtccaaagg gtgccaatga	480
ggtgctcctt	gtgatcgagg cgtacaagac gctgagagac agagcacctt accctgctag	540
cttcatgctc	tcgcagctcg ccggcagcta cgcccttgcgt ctcttcgaca agtccaccc	600
cacccttc	gtggcctctg atccggaggg aaaggtgtct ctgtattggg ggattactgc	660
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gtgctatgag	aacccaaagc acaaggtgac cgcaatgttcc gccaaagaag aagaaatttg	840
cgggtcaacc	ttcaagggtgg aaggatctac aatcctcaca ggcgtgcatt aggagattt	900
tggatctcca	gctgttgggg ggaaaggaaa actctgttagt ctgtaaatgg tgggttaaca	960
tagtcgcaag	gcatgcacat gatctgtatc atcatctcc gtaaagacta gtgtgcacag	1020
catggtttagc	acttttctgtt gttttttgtt gtgtatgtatc tttatcttat gttgttcagg	1080
cgaaaaatgaa	tttcaggcccc agctgtgtca gttgaatgta gttgtgttc	1130

<210> SEQ ID NO 12  
<211> LENGTH: 1653  
<212> TYPE: DNA  
<213> ORGANISM: Oryza sativa  
<220> FEATURE:  
<221> NAME/KEY: misc\_feature  
<223> OTHER INFORMATION: LOCUS: AK065009 1653 bp mRNA linear PLN  
04-DEC-2008; DEFINITION: Oryza sativa Japonica Group cDNA  
clone:J013001E24, full insert sequence.; ACCESSION: AK065009

<400> SEQUENCE: 12

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gccccatgtgt tggtgcgtcg tcggcggtct tggtcttggt tggtcgaaaa ctgtgattac	180
aaagagagag gctcgagtgg tgaattttca cgggaaaccc tagegaggcg gcggatctcg	240
ccgtgtgccc ccctccccc cacgagcacc ccgtgtgttc acggcggtgg agtggatgga	300
acccgagggtg ccgcggccatc gcgatgttgg ctgggttata tcggtagttg catccatggc	360
tcaccatgt agtactactc cctcccccgtc caaaatgtta ttgtacttc ctccgttca	420
taagggtata agttcatttc tggtggagga atggatggcg cggtgggtgg gggtgctgtc	480
tctgtgagggg gtttctgtcc acgatgaagg cttcgtttt aacgcataaca cttgatatga	540
gcccccttta tgatggatt ttgtatgtat gtatgggtt ggcgagctga attcctgttc	600
ttgcaacatc tgatggatgg tgcgatgtat ttgcttggat taccgtttt tcaatgtatca	660
taggaaatata gtttatcca ttatgtctg aagatgtca ttaatgttgg ggtgtcaact	720
gtttaaataa tactccgttag tactgtctgt agggaaattat aactattata tactaatagt	780
agctgttaag ttgatcgatc attatgtat gtcactgtc tatatgtat tccgtgtgt	840
gtttcagtag gaaattatga atgcttcaa ttccctgtga acatgaaatt cagttacata	900
actgcattat cggtgattac ctaagttacc atttcatagt agttatgtt gggtgtcaaa	960
gtttatccag ggaagtacat cttatcatt cattcatgtat ttgatagact ttgtatatg	1020
aaattgtcat tctgtatgtt gtcaatctgtt attgccattt tgccatccat gatgccccta	1080
cttcagttgg ttgttcaac aatgtctatg ctggcattct tgaatcaccc aagcaatgaa	1140
tttaaatcaa cttaatttgc tttgtatgtt tttgtatgtt tgcttctgg ttccagttc	1200
tgttcaccct tttgtatgtt gagagaaggc tactgtgttg tttgtatgtt cagtttctta	1260
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ctgtttgtat tttgtatgtt ggaagaagaa ggcgtatggc aggctgttgg ggaaggcccg	1380
aaagatgagg cagatgtca agtggcgaa ggagatgggg gatcatgggtt atgggtgtc	1440
gcagtgatcc atgtttcggt ctggcaagct cttgtatgtt tcaatgttgg tggcggtgg	1500
ccaccattcg actgttgaat ggcttattta cttgtgtgtt tatcatctt ttttgtatgtt	1560
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gtatgtatgtt cttatgttta cttaccgttg agt	1653

<210> SEQ ID NO 13  
 <211> LENGTH: 805  
 <212> TYPE: DNA  
 <213> ORGANISM: Oryza sativa  
 <220> FEATURE:  
 <221> NAME/KEY: misc\_feature  
 <223> OTHER INFORMATION: LOCUS: AF017360 805 bp mRNA linear PLN  
     28-MAY-1999; DEFINITION: Oryza sativa lipid transfer protein LPT  
     III mRNA, complete cds.; ACCESSION: AF017360

<400> SEQUENCE: 13

gcatcagcaa ccaattcgca ccgatcgatc gatcgatcga tccagcaact agatcaacag	60
tactagctat gaagcatggc ccgtgcacag ttgggtttgg tcggccgttgg ggcagctctg	120
ctccctcgcc ccccgacgc cggcggtggcc atcacctcgcc gccagggttcaaa ctccggccgt	180
ggccctgtcc tgacctacgc ccggggcgcc gccggccgttgg cggcggttgg ctgcaacggc	240
gtgaggagcc tcaaggcgcc agcaaggcacc accgcagacc ggccgaccgc ctgcaactgc	300

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ctcaagaacg	cggccgcgg	catcaagggg	ctcaacgccg	gcaacgcgc	cagcatcccc	360
tccaagtgcg	gcgtcagcgt	ccctacacca	tcagcgcttc	catcgactgc	tccagggtga	420
gctgagccat	cgatcagaga	gacggatcat	atatatacag	cagcgcgcgg	gttgcacact	480
atgtagattct	gcctgggtgc	atgtgtggac	ccgaattctg	tatccagtac	tagtagtata	540
atatctgtat	tctggaataa	aagatgagct	agctaaggtc	tcgatcaatc	accatgcatg	600
catgtgtgt	catccatgg	tgatcgccgc	gtcacgtag	ctagcttct	tcttttgt	660
tgttctgtctc	gtacgttttgc	ctcccttctg	aggggtacgt	gtaccagaga	gagctagaga	720
ttcttacatgc	atgtactgca	actccttgc	ctacgtgttt	tgtttggat	tattacacat	780
acatatactt	cttttcgtt	cattt				805

<210> SEQ ID NO 14  
<211> LENGTH: 1130  
<212> TYPE: DNA  
<213> ORGANISM: Oryza sativa  
<220> FEATURE:  
<221> NAME/KEY: misc\_feature  
<223> OTHER INFORMATION: LOCUS: AK071205 1130 bp mRNA linear PLN  
04-DEC-2008; DEFINITION: Oryza sativa Japonica Group cDNA  
clone:J023085D08, full insert sequence.; ACCESSION: AK071205

<400> SEQUENCE: 14

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gcaaaaatcc	gatcctcctc	ctgcctctgt	cgctcgctcg	ctgcgcgcac		180
ttggccccgt	cccgtccgtt	cccgctcgcc	aggcatagaa	agatgggtct	ctgggtcttc	240
ggctatggct	ccctgtatctg	gaaccccgga	tgcacttcg	atgagaagat	cctgggtttc	300
gtcaagggt	acaaacgcac	cttcaatctt	gcttcatttgc	accatcgccc	cacgcggag	360
catcctgcga	ggacctgcac	acttgagtcc	gacgaggaa	ccatatgtcg	ggggatttgc	420
tactgtgtca	aaggggggct	taaaaaggag	caagaagcaa	tgaagtactt	ggaaaagaaga	480
gagtgtgagt	atgaccagaa	gatctctgt	gatttctaca	aggaaggaga	ctctttgaag	540
ccagctgtga	cagggttact	agtctttgt	tccactcctg	atccagtagg	caacaaatac	600
tatcttgggc	ctgctccctt	ggaggacatg	gcaaggcaaa	tgcctacagc	caatggcccc	660
aatgggata	acagggatta	cctgttctca	atggagaagg	cattgtccaa	catatgccat	720
gaagatgatt	caatcatcga	gctagctaac	gagggtcagg	agggtgttgc	ccggccgaaag	780
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tccttggcac	actggggAAC	cagaatgtgc	tgatTTCTT	tgcacccgc	tgcggcaatg	1020
ctgttaactt	catttgcata	atactagac	cgagactgg	gtgctgtgt	ttcccttgac	1080
acactaaata	ataaaaggat	tccatgagct	tttcactctt	gagctgttgc		1130

<210> SEQ ID NO 15  
<211> LENGTH: 779  
<212> TYPE: DNA  
<213> ORGANISM: Oryza sativa  
<220> FEATURE:  
<221> NAME/KEY: misc\_feature  
<223> OTHER INFORMATION: LOCUS: AK059164 779 bp mRNA linear PLN  
04-DEC-2008; DEFINITION: Oryza sativa Japonica Group cDNA

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clone:001-023-D11, full insert sequence.; ACCESSION: AK059164

<400> SEQUENCE: 15

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accgctgctt	cgtcgccggc	ctcgccctgg	ccaccgacga	ccgctccctc	gagggccct	180
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tatcggtgtcc	agtggctctc	tcgagtcgag	aaggccctcta	tccatccatc	cagtgttagg	660
tgttcttcgt	ccgtgtatgtt	accatgaatt	gagttcgctt	tggttatgtt	gtttgaactg	720
cttgggtcta	tctatcgaa	tgaaatgaaa	tagaaaacaa	ggagaaaaaa	aagagttcg	779

&lt;210&gt; SEQ\_ID NO 16

&lt;211&gt; LENGTH: 1667

&lt;212&gt; TYPE: DNA

&lt;213&gt; ORGANISM: Oryza sativa

&lt;220&gt; FEATURE:

&lt;221&gt; NAME/KEY: misc\_feature

<223> OTHER INFORMATION: LOCUS: AK107238 1667 bp mRNA linear PLN  
04-DEC-2008; DEFINITION: Oryza sativa Japonica Group cDNA  
clone:002-125-F02, full insert sequence.; ACCESSION: AK107238

&lt;400&gt; SEQUENCE: 16

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gtccgaacgt	aaatccatgg	cacaaccctc	ggcaaggagg	tttttagggag	tgttagattt	180
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actttgtat	gaagaatgaa	caattccagt	gcacaggatc	atttgcatac	cgacgtgtca	300
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tccaaggagg	aggttctatg	ggattaactt	tccccggctg	cccagcaacc	taccaacaac	420
aattccaaca	attcttgcc	gaaggccaaa	gccagagcca	aaaatttagg	gtgagcacc	480
aaaagatcca	ccaattttaga	caaggagata	tcgttgact	gccagctggt	gttgcgcatt	540
gtttctaca	tgaaggcgat	gcaccagtg	ttgctctata	tgtcttcgac	ttaaacaaca	600
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accaagttca	gtacagtgaa	gaacaacaac	catctacgct	ttgcaacggt	ttagatgaga	960
acttctgcac	aatcaaggca	aggttgaaca	tcaaaaatcc	tagccatgt	gataactaca	1020
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gttaaccttgg	aaagacggta	ttcaatggcg	ttctacgtcc	aggtcaatttgc	ctgatcattc	1260
cgcaacacta	cgttgtcttg	aagaaagcag	agcatgaagg	atgccaatac	atttcatca	1320
agaccaatgc	aaactccatg	gtgagccacc	ttgcaggaa	gaactcaata	ttccgtgcca	1380
tgccagtgg	tgtgatcgct	aatgcttacc	gcatacgtgg	ggagcaagca	cgaagcctta	1440
agaataatag	gggagaagag	ctcggtgct	tcactcttag	atatcaacaa	cagacctacc	1500
taggcttc	aatatgatcg	gagaacgagg	cctcagagtt	atgtactaat	gaaatagttat	1560
aggtgtatca	aaaaaaaata	aatgccaca	agtatgtgaa	actttgtggc	ggttctgttc	1620
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<210> SEQ\_ID NO 17  
<211> LENGTH: 617  
<212> TYPE: DNA  
<213> ORGANISM: Oryza sativa  
<220> FEATURE:  
<221> NAME/KEY: misc\_feature  
<223> OTHER INFORMATION: LOCUS: AB016505 617 bp mRNA linear PLN  
09-JAN-1999; DEFINITION: Oryza sativa mRNA for prolamin, complete  
cds, clone:lambda RM9.; ACCESSION: AB016505

&lt;400&gt; SEQUENCE: 17

gaaaaaggat	aagaactaga	aaccaccac	aatgaagatc	atttttttct	ttgctctct	60
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cgttaaggcg	cagtgcagca	cagtggcaac	cccccttctt	caatcacccg	tgtttcaact	240
gagaaaactgc	caagtcatgc	agcagcagt	ctgccaacag	ctcaggatga	tcgcacaaca	300
gtctcactgc	cagggcatta	gcagtgttca	ggctattgt	cagcagctac	ggctacaaca	360
gtttgctagc	gtctacttcg	atcagagtca	agctcaagcc	caagctatgt	tggccctaaa	420
catgcccgtca	atatgcggta	tctacccaag	ctacaacact	gtccctgt	gcattccac	480
cgtcggttgt	atctggatt	gaattgtgc	agtatagtag	tacaggagag	aaaaataaaag	540
tcatgcac	tcgtgtgtga	caagttgaaa	catcggtgt	atacaaatct	gaataaaaaat	600
gtcatgcac	tttaaac					617

<210> SEQ\_ID NO 18  
<211> LENGTH: 1460  
<212> TYPE: DNA  
<213> ORGANISM: Oryza sativa  
<220> FEATURE:  
<221> NAME/KEY: misc\_feature  
<223> OTHER INFORMATION: LOCUS: AK099086 1460 bp mRNA linear PLN  
04-DEC-2008; DEFINITION: Oryza sativa Japonica Group cDNA  
clone:J023009G10, full insert sequence.; ACCESSION: AK099086

&lt;400&gt; SEQUENCE: 18

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cccgctccgg	tgatcgatc	cccacagtt	tgcgtttca	ctccctctca	tctccattcg	180
tttttgagtt	ctcggttgc	tccgccttc	tttcaactat	gggcaagatt	aagatcgaa	240
tcaatgggtt	cggccgcac	ggcaggctgg	tggccagggt	ggcgctgcag	agcgaggatg	300
tcgagctcg	tgccgtcaac	gatccctca	tcaccaccga	gtacatgaca	tacatgttca	360

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atgtatgcacac cgttccacggc cagttggaaac atcatgaggat caagggtcaag gactccaaga 420  
ccctcatctt tggcacgaaa gaggttgccc tggttcggctg caggaaccct gaggagatcc 480  
catgggctgc ggctgggtct gaatacgttg tttagtctac tggtgtttc accgacaagg 540  
acaaggcaggc agctcaacttg aagggtggtg ccaagaaggt cgtcatttct gctcccgaea 600  
aagacgcccc catgttcgtt gttgggtgtca acgagaagga gtacaagtct gacgtaaca 660  
ttgtctccaa cgcttagttgc accaccaact gcctggctcc tctcgccaaag gtcataatg 720  
acagattgg catcggttagg gggtttagtga ccactgtcca tgccatcaact gctacccaga 780  
agactgttga tggggccctcg atgaaggact ggagaggtgg aagggtctgct agcttcaaca 840  
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gtgggtgggtt gggccggc atggctcata ttttgggtct aatttttttgc cgcttaatct 1440  
aaatcqaatq qttqcttcgc 1460

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<210> SEQ ID NO 19
<211> LENGTH: 942
<212> TYPE: DNA
<213> ORGANISM: Oryza sativa
<220> FEATURE:
<221> NAME/KEY: misc_feature
<223> OTHER INFORMATION: LOCUS: AY166458 942 bp mRNA linear PLN
    23-MAY-2006; DEFINITION: Oryza sativa (japonica cultivar-group)
    alpha-amylase/subtilisin inhibitor mRNA, complete cds.; ACCESSION:
    AY166458
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<400> SEQUENCE: 19

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cggtgtacga cacggagggc caegagactga gggccgacgg gagctactac gtcctccgg 180
ctagccccgg ccacggagggg ggccctcaega tggccggcccg cgtgtcccccc tgcccgctcc 240
tcgtggggca ggagacggac gagccggca aaaaaaaaaaa cgtgcgtttc accccgtggg 300
ggggccggcc ggccggggag gacaggacca tccggctctc gaccggacgtc cgccatccgct 360
tcaacggccgc gacgatctgc gtgcagtcga ccggatggca tggccggcgtc gageccgtca 420
cgggggccgc gcgcgtggtg acggggccgt tgatggggcc gagccggac ggggggggaga 480
acgcgttccg cgtggagaag tacgggggtg ggtacaagct ggtgtcgatgc agggactcg 540
ggccaggaccc gggccgtgtca agggacggcg cgccgggggtg gctggccgcg agccagccgc 600
ctcacgtcgat ggtttcaag aaggccagggc caagccacc agagtaaacg agggggggggg 660
aaaaaaaaaaaa ggtcactcat gcgttgcgtg gtgcgggtgtt gctctgttag cgtgtatagg 720
tagtaggggtt cgtgcgttgcg tgccatgtc ctgggttaat aatgtgtgaa gaggcttagca 780

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ggtattgctc cgtggttctg ttctcatgtt cgtgtcatgt ctgttagtcg tgataactcg	840
tgagcactga gcaagtggc aggactgcct ctggctgtct ggcccgccg gacttgtaa	900
gtccaaatgg aatgaacgaa ttgtgcgggaa taaaaaaaaaa aa	942
<210> SEQ ID NO 20	
<211> LENGTH: 1588	
<212> TYPE: DNA	
<213> ORGANISM: Oryza sativa	
<220> FEATURE:	
<221> NAME/KEY: misc_feature	
<223> OTHER INFORMATION: LOCUS: AY987390 1588 bp mRNA linear PLN 19-APR-2005; DEFINITION: Oryza sativa (japonica cultivar-group) glutelin precursor (GluB7) mRNA, complete cds.; ACCESSION: AY987390	
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tctatggccc aactatTTAA tcAACAGCACA aACCCATGGC atAGTCTCG gCAAGGAAGT	180
tttaaggagt gtaggTTTGA tagactacaa gcATTGAGC cACTTGAAA agTAAGGTCA	240
gaagctgggg tgactgagta cttcgatgag aagaatgaat tattccaatg cacaggtact	300
tttgcattcc gacgtgtcat tcaacctcaa ggcctttgg tacctcgata tagcaatact	360
cctggcctgg tctacatcat tcaaggggagg ggttctatgg gtttaacctt ccccggttgc	420
ccagcgactt atcagcaaca attccaacaa tttcgtctc aaggccaaag tcaaagccaa	480
aagtTTAGGG atgagcatca aaagattcat caatttagac aaggagatgt tggactc	540
ccagccggtg ttgcacattt gttctacaat gatggtgatg catcggttgt tgccatatat	600
gtttatgaca taaacaacag tgcaaatcaa cttgaaccaa ggcaaaagga gttcctatta	660
gctggtaaca acaatagggt tcaacaagta tatggcagct caattgagca acactctagc	720
caaaacatata tcaacggatt cggtactgag ctactaagt aggctttagg catcaacaca	780
gtagcagcaa agaggctgca gagccaaaat gatcagagag gagagatcgt acatgtgaaa	840
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taccaagagg ttcaatatacg tgaacaacaa caaacatctt cccgatggaa cggattggag	960
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ctcatccaaa tgagtgtac cagagtaaac ctataccaga atgctattct ctcaccattc	1140
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gccttaccag ttgatgtggt cgctaattgt taccgcattt cacggagca agccccaa	1440
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<210> SEQ ID NO 21	
<211> LENGTH: 1764	
<212> TYPE: DNA	
<213> ORGANISM: Oryza sativa	

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<220> FEATURE:  
 <221> NAME/KEY: misc\_feature  
 <223> OTHER INFORMATION: LOCUS: AK107314 1764 bp mRNA linear PLN  
 04-DEC-2008; DEFINITION: Oryza sativa Japonica Group cDNA  
 clone:002-126-D11, full insert sequence.; ACCESSION: AK107314

&lt;400&gt; SEQUENCE: 21

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agagcactag tcaatggcag agttctcg tc gtggaaatcc gagaggatgt agatttgata	180
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tgcgtgtctc taatgagttt tttcaatgtta ccggagttatc tggtgtccgc cgagtattt	300
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aaggggaggg tataacaggg ccgactttcc caggctgtcc tgagacctac cagcagcagt	420
tccaacaatc agggcaagcc caattgaccg aaagtcaaag ccaaagccat aagttcaagg	480
atgaacatca aagatcac cggttcagac aaggagatgt tatcgctgtg cctgctgtg	540
tagctcattt gtcgtacaat gatggtaag tgccgggtt tgccatatat gtcactgata	600
tcaacaacgg tgctaatcaa cttgaccctc gacagaggga ttcttggta gctggaaata	660
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gcttttagcac tgaactgctt agcgaggctt ttggcataag caaccaagtt gcaaggcagc	780
tccagtgtaa aatgaccaa agaggagaaa ttgtccgcgt tgaacgcggg ctcagttgc	840
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atcaagaagg aggatatacg caaagtcaat atgggagtg ctgccttaac gggttggat	960
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tcaacaacaa tggaaagacg gtgttctacg gagagcttcg tgcgtggacag ctacttattt	1260
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aagacgttta taatgtggcg gaatcccttt aagttggtaa tgccgataaa gaataactaa	1560
ataaataaat aaataaattt caagcaattt cggtgtgtc atgtactgtt aaagtttctt	1620
ataatatacg ttctgaatgc taaggacatc cctcaagatg gtctttctat ttttgttcc	1680
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<210> SEQ\_ID NO 22  
 <211> LENGTH: 1637  
 <212> TYPE: DNA  
 <213> ORGANISM: Oryza sativa  
 <220> FEATURE:  
 <221> NAME/KEY: misc\_feature  
 <223> OTHER INFORMATION: LOCUS: X15833 1637 bp mRNA linear PLN  
 18-APR-2005; DEFINITION: Rice mRNA for glutelin.; ACCESSION:  
 X15833

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<400> SEQUENCE: 22  
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caaaccatg gcatagtcctt cggeaaggaa gtttttaggaa gtgttagattt gatagactac 180  
aagcatttga accaattcgg aaagttaggtt cagaagctgg ggtgactgag tacttcgtat 240  
agaagaatga attattccag tgcacggta cttttgtat ccgcacgtgtc attcagccctc 300  
aaggccctttt ggtacttcga tacacaaaata ttccctggcgtt ggtctacatc atccaaggaa 360  
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atgtatgttgcaccatc aacatccatc tggccgttgc ttttgcaccatc 1560  
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aagggttacaaacttctt 1637

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<210> SEQ ID NO 23
<211> LENGTH: 887
<212> TYPE: DNA
<213> ORGANISM: Oryza sativa
<220> FEATURE:
<221> NAME/KEY: misc_feature
<223> OTHER INFORMATION: LOCUS: AK061894 887 bp mRNA linear PLN
        04-DEC-2008; DEFINITION: Oryza sativa Japonica Group cDNA
        clone:001-041-G07, full insert sequence.; ACCESSION: AK061894
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catgaccgtc ggcggagctc cggcgggcg gatcgtgtatc gagctgtacg cgaaggacgt 180  
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gttcgecgac gaggtgttca agttcaagca	cgacagecccc ggcatectgt ccatggcgaa	420
cgcggggccc aacactaacg ggtcccaagtt	cttcatctgc accgtgcct gcagctggct	480
ggacggaaag cacgtcgtgt	tcggccgegt cgtcgaggc atggacgtcg tcaaggccat	540
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tcgtgttggt gagatgagat cggccatgtt	ttgggtggat taggcggagt tcttggatcg	780
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<210> SEQ\_ID NO 24  
<211> LENGTH: 1650  
<212> TYPE: DNA  
<213> ORGANISM: Oryza sativa  
<220> FEATURE:  
<221> NAME/KEY: misc\_feature  
<223> OTHER INFORMATION: LOCUS: AK107343; 1650 bp mRNA linear PLN  
04-DEC-2008; DEFINITION: Oryza sativa Japonica Group cDNA  
clone:002-126-G05, full insert sequence.; ACCESSION: AK107343

<400> SEQUENCE: 24

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ttaaatccca gcacaaaccc atggcatagt	cctcggcaag gaagtttag ggagtgtaga	180
tttgatagac tacaagcatt tgaaccactt	cgaaaaagtga ggtcggaaagc tgggtgtact	240
gagtaactcg atgagaagaa tgaattattc	cagtgcacgg gtactttgt gatccgacgt	300
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caacaattcc aacaattttc atctcaaggc	caaagtcaaa gccaatgtt tagagatgag	480
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cattggttct acaatgtatgg	tgtgcacct attgtgcgg tatatgttta tgacgttaac	600
aacaacgcca atcagcttga acctaggcaa	aggaggttc tattagccgg caacaacaat	660
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agtaactttg gatagactgt	gtttgatgtt gtccttcgccc caggacaattt attgtatcatt	1260
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ccaggattat cgaatgagtc cgaaagcgag	acctcagagt aatgtatgt agaactagta	1560
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<210> SEQ ID NO 25  
<211> LENGTH: 1556  
<212> TYPE: DNA  
<213> ORGANISM: Oryza sativa  
<220> FEATURE:  
<221> NAME/KEY: misc\_feature  
<223> OTHER INFORMATION: LOCUS: AK063995 1556 bp mRNA linear PLN  
04-DEC-2008; DEFINITION: Oryza sativa Japonica Group cDNA  
clone:001-124-E11, full insert sequence.; ACCESSION: AK063995

<400> SEQUENCE: 25

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tgaggaggag aacatggcg	cgtggctgtt ggccaaagaac accctaaga tcatgcctt	180
caagctcccg ccagttgggc cttatgtatgt	ccgtgtccgg atgaaggcag tgggcattctg	240
cgccagcgcac gtgcactacc tcagggagat	gcccattgcg catttcgtgg tgaaggagcc	300
gatgggtatc gggcacgagt gcgcggcggt	gatagaggag gtccgcggcg gcgtgacgca	360
cctcgcgcgtc ggcgaccgcg tggegctega	gccccgcatac agctgtggc gctgcaggca	420
ctgcaaggcgc ggccgcatac acctctgcga	ggacatgaag ttcttcgcac cccctccgt	480
ccacggatcc ctgcacaacc agatcgtca	ccctgggtat ctgtgtttca agctgcggga	540
gaacgtgagc ctggaggaag gcgcacatgt	cgagccgtcg agcgtggcg tgcaegcgt	600
cgcgcgcgc gacgtcgccc cggagacggg	gggtgctgatc atgggcggcg ggccgatcg	660
cctggtcacc ctgctgggg cgcgcgcgtt	cgccgcgcac cgctgttgta tcgtggacgt	720
ggacgaacac cgcctctccg tggcccgatc	cctcggegcg gacgcgcgcg tgagggtgtc	780
ggcgcgcgcg gaggacgtcg	cgccggatcggt ggaacggatc agggcggcga tgggggggaa	840
catcgacgtg agcctggact gcgcgggtt	cagcaagacg gtggcgacgg cgctggaggc	900
gacgcgcgcg ggccggaaagg tgcgttgt	cggtatgggg cacaacgaga tgacgggtcc	960
gtgtacgtcg gcggcgatca gggaggtgga	cgtgggtgggg atattccgtt acaaggacac	1020
gtggccgctc tgcacatcgat	tcctccgcag cggcaagatc gacgtgaagc cgctcatcac	1080
ccaccgatcc gggttctcg	aggaggacgt ggaggaggcg ttcgaggatca gcgcgcgtgg	1140
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agcaaaaatat gtgacgggtt	tggAACGTAT gtcgatcgga caaaaaaatg tgtgttgct	1260
ttgtctggttc ccgtgtctgt	ctgtctctac atgaataaaa ggccggcaacg ttctgtcctc	1320
atggactaag cgtgtgcatt	gtctgtttca attcattcgta tggtgttga acactagggg	1380
tgggggggg gaaatcgaaa	ctcgcagctc ggatcggtt gtagcaagtt cggcttgat	1440
cagcttggac tcgtaatctt	aatgagtcga gctgagtag ccttttaact catgagctc	1500
tcgtttaact cgtagtaca	agatataacc aaaccttcgt aaacttggtg cttttt	1556

&lt;210&gt; SEQ ID NO 26

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<211> LENGTH: 831
<212> TYPE: DNA
<213> ORGANISM: Oryza sativa
<220> FEATURE:
<221> NAME/KEY: misc_feature
<223> OTHER INFORMATION: LOCUS: AK108210 831 bp mRNA linear PLN
  04-DEC-2008; DEFINITION: Oryza sativa Japonica Group cDNA
  clone:002-140-D09, full insert sequence.; ACCESSION: AK108210

<400> SEQUENCE: 26

accatttca ctcactcaact cacactgaac tcctatggcc tccgcccgtcg tagccaagcc      60
accggccgccc gtcgggcga gcccggcgg atgcttcgtc ttctgttaggc gcgcggccgc      120
cgccgcgcgc ggcgtcaacg cggcgatcgc tgccgtcgat gaccacccgc cagttccggc      180
cgccgcgcgc gccgecatgg acgacgtcg tggccgcgtc ggcggccgtc cgccgcgcag      240
cgccgcgcgc atcatggagg ggacgcacaa gcagatcgc tcggggggcg cctcgccgg      300
ctactgcacc gtgcgttgtt gcagcatctg caccggaaat aaccctgtcg ccattgcga      360
gttcttgctg tgctgcaacc tctggggcgat ccctctcgcc ggcgcgcgc gcttcatcta      420
cattggagag aaggcggtct gcaaggagga gtgcagggtcg aggtacgtgg tggaggaggc      480
gtgtgcgcgag gcgaggaggagg agaagcgccg cgcgcgcgc ggcgcgcgc cgccgcgg      540
gaagaagaag gagggggcgcc cgccgcggaa gggggggggag gactgcagggg agggggagcat      600
cttcttcatc tgcgcgcgacg acctgtgaag aatgtgtatgg atatgtatgc tcatgcgtgc      660
attgcatggc ggtatataat cggagcttgg ctgtatataat aatcgatctc caccataata      720
taatacaata gtaataact tagtgactg taatgaggat gaataaaaggg gatcaaata      780
aggccaccat gcatggctag tcgatataata tctgagtatt ttgtgttgc c      831

<210> SEQ ID NO 27
<211> LENGTH: 1344
<212> TYPE: DNA
<213> ORGANISM: Oryza sativa
<220> FEATURE:
<221> NAME/KEY: misc_feature
<223> OTHER INFORMATION: LOCUS: AK242579 1344 bp mRNA linear PLN
  04-DEC-2008; DEFINITION: Oryza sativa Japonica Group cDNA,
  clone:J090009I07, full insert sequence.; ACCESSION: AK242579

<400> SEQUENCE: 27

ttggatcatc gaatctcgca cggccactgg aacttgccaa cccgagaatg acggcattcc      60
aggacaacac catggccaat gggacggcac cgccgatcag tattgcgcgc cgccatcg      120
acactgaatc cccccacaatc ttgcagatga acggcgatgt gacatggaa ccaagcgatca      180
ggacggtcac cgaaattgca ggcaggatgg tagcaggcgcc ccagcaggcg tagccgagcg      240
atctcagcat gctgtccgg ccgacggaaa cgccgggtgac aacaagcgcg gtgtatggagg      300
cgagcatcag gccgcagagc ggcgggttgg cgccgtcgac ggccttgaag gggagaacg      360
cgatgagcgt cccggcgaag cacgggaaga tggcggttggc caaggccgg tccaccggcc      420
ggaagagctg ggacaccagc gagccgatgc cggcgacgc ggcgacgagc agggagaagc      480
tgagggccgc gtagacgacg gccacgaccc ccccgagcgt cgcccccaag gtgtcgacg      540
caagtccctgt gaagctgacc tcgtcgacgc cgccgtctc catcgccggc aagctgagct      600
ccgcgacgag gacgtggag gacaccacgt agacccacga gggcacgatc gccgcgtgg      660
acggggacccg cccggagcgg atgggtggcg acggcagccg cagcatcccc ggccccacccg      720
cggttcgac gatgaggctc accggccgccc agaagctctt cttctccctt ccacccgata      780
ccccgtctc gacgagcggc gccgggtgatt cttcttcggc ggcgtcgccg tcggcgtagt      840

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cgcgtccgcca	ccggcattgc	cgcctcgcg	acagccagga	cagccgcaac	ggctgcggct	900
ggggcgccgc	ggggttcttg	gaagctatgt	ggggaggcg	acgcgttgcgt	ggctccggga	960
ggccgaggcg	ggggggcg	cgagccggag	agggggcaggt	tgcggcgag	gagggcgtgg	1020
cgaggtggag	ggggctggag	agtaacatgg	tcaacggtaa	attttgttag	ggtcaaagga	1080
ggcggtgaa	ttctggccct	ctttaccat	tcgaggccc	gttttcaac	atacgagtt	1140
ctaaaacatt	ttggatttggg	aatttgaaga	atgaataata	atacgtatgt	caaactgagc	1200
ggattcatgt	aaaaaaatac	tgtaaaattt	ctgcgtctca	atggagccctc	agcaaggcct	1260
tgaaagattt	gggatttagag	aaattaatat	ggctgaattt	acgtttccat	tgaaacggaa	1320
gaatttttagt	gaattttcat	tatc				1344

<210> SEQ ID NO 28  
<211> LENGTH: 2078  
<212> TYPE: DNA  
<213> ORGANISM: Oryza sativa  
<220> FEATURE:  
<221> NAME/KEY: misc\_feature  
<223> OTHER INFORMATION: LOCUS: AK065456 2078 bp mRNA linear PLN  
04-DEC-2008; DEFINITION: Oryza sativa Japonica Group cDNA  
clone:J013025D11, full insert sequence.; ACCESSION: AK065456

<400> SEQUENCE: 28						
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caggaaatcc	ggctccggtc	cctgtgccgc	tccccccggc	tctccttgcc	tcccttgctc	120
ctccgatcga	ggctgaagcg	gcttcaggat	atacatTTT	gtggaaatgg	ggaattcttg	180
ccaaaatgga	acctatggga	acaattacca	gaacagcaac	cggtttcaga	atgaccgttt	240
tgtttctcg	tacgttgcgt	ggaatgatac	tgaggattgc	tactcggtct	cgtcaaggc	300
cagcttagcg	ggtgctctgc	ggcaaggcgt	gaacctaaag	tcccctgtcc	ttggatacaa	360
gactccaaat	gtaggggagc	tctatactct	tggccggggag	cttggacagg	gacagttcg	420
gaaaacatac	ctctgcactg	agattagcac	agggtgtcaa	tatgcatac	agactatctt	480
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gcaccatctt	tcggccaga	agaatatagt	gacaatcaag	gacacatatg	aggatgagca	600
ggccgtgcac	atcgcatgg	agctctgtgc	aggtggtag	ctttcagca	agattcagaa	660
gcgaggtcat	tacagtgaac	ggaaggcgtc	agagctata	aaaattatag	ttggcatcat	720
agaaacatgc	cattcacatg	gagtgtgc	ccggatctc	aagccagaaa	atttcctctt	780
actggacgc	gacgatgaat	tctcggttaa	agcaattgac	tttggcttat	ctgtgttctt	840
cagaccaggt	caggTTTca	gagaggtgt	gggaagtcc	tattatattg	ctccagaggt	900
atggagaag	cggttatggac	cagaggctga	tatatggact	gtggagtg	ttctctatgt	960
attgtctact	gggtgttctc	cattttgggc	agatacaca	agccggatata	atgaaaaagt	1020
actggatgga	cgtattgtt	ttaaatcaa	ccggtgccc	aggatatactg	acagtgc	1080
ggatcttata	aaaaagatgc	tctgcctta	tccatcagag	cgtttggaaag	cccatgaagt	1140
gctaaagcat	ccatggatata	gtgataatgg	agtggctact	aatcgac	tggatccaag	1200
tgtacttcct	cgtctcaagc	aatttctgc	aatgaatagg	ttaaagaaat	tgtctctcca	1260
gattattgt	gagcgtcttt	cagaagagga	gattgttggg	ttaagagaaa	tgttcaaggc	1320
tatggacacc	aaaaacagaa	gtgtggttac	atttgggtgag	cttaaggac	tgaaaagata	1380
cagctcagtg	ttcaaggata	ctgaaattaa	cgacttaatg	gaagcagctg	atgacaccac	1440

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ctctaccatc aactggaaag agtttattgc tgcagcagtgc tctctaata aaatagaacg 1500
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agttgacaag cttcaaaagg cttgcatttgc acgttaacatg gaagatactt tccttgaaga 1620
gtgattctg gagggttgc acaaacaatga tggtaaatttgc gattatgtcg aatttgcac 1680
aatgatgcaa agcaacaact ttggacttgg gtggcaaaccg gtggaaagca gcctgaatgt 1740
agccctgagg gaggcaccgg aagtataactg aactcctgtc gcctggcacc cccaagaatt 1800
cagtttgc tccatgttt tatcacaaca catagttgtg gatttggag 1860
aagagagcca agacttagatg gtttatcagt aatcaccctt tagcagtgtt ggaagaagac 1920
tctcttagta octaatagca tataaatgtt tggtaaatttgc acatcgata tggatgtt 1980
tcataagttc gttcggttgc tggtttttttgc taccctgtc atctaataat 2040
aatatcaata ataccgaaag aacccttgc gtttgc 2078

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<210> SEQ ID NO 29
<211> LENGTH: 729
<212> TYPE: DNA
<213> ORGANISM: Oryza sativa
<220> FEATURE:
<221> NAME/KEY: misc_feature
<223> OTHER INFORMATION: LOCUS: AK107633 729 bp mRNA linear PLN
    04-DEC-2008; DEFINITION: Oryza sativa Japonica Group cDNA
    clone:002-131-E06, full insert sequence.; ACCESSION: AK107633

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<400> SEQUENCE: 29
aaaaattaaa ttttctaaat tacaagcaaa tcaaagctca tcgaagtata gctatggcat 60
tggcatcaga caagttcgcc ctctccgcca tcgtgctcg cgtcctcacc gtcggcggcag 120
cggcggcggg ctacggcggc tacggcgacg tcggcgact ctggcgctg gggaggcgg 180
tgtccccggaa cccggtccca tcgtgccggaa actacatcgcc ggggtggtgc gccgtcgccg 240
ggggccgcct ggactccggc aagcagccgc cgccgcgcgt cctggagccg tgctgcccgg 300
agtcgcgcgc agtgcgcgtg cagtgcgcgt gcgcgcgtt gagcgtgtgc gtgcgtggcg 360
tggtcacggaa ggagggcgac cgctcgccgc ggatgatctc gcagcacgcg gcccgggt 420
gagaegccgc gacgatcgcc gggatggcga ggcgcgtgac ggactacggc cggtgcacc 480
tgcagcacac tggttcttt ggctgccccaa tggatgggggg tggatggat taacttcctt 540
agtaattaaat taattaggcc tttgttaat taattattta attagttatc cgggttactg 600
gataattaaat tatcgatatt tggatgtgc atctatcatg tttggatgtc gctttctccg 660
tgaatgtgat gataataata atcagaagaa ataaataaga agagttggat tcatcagctt 720
tccagttatc 729

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<210> SEQ ID NO 30
<211> LENGTH: 1507
<212> TYPE: DNA
<213> ORGANISM: Oryza sativa
<220> FEATURE:
<221> NAME/KEY: misc_feature
<223> OTHER INFORMATION: LOCUS: AY196923 1507 bp mRNA linear PLN
    02-AUG-2005; DEFINITION: Oryza sativa (japonica cultivar-group)
    glutelin precursor, mRNA, complete cds.; ACCESSION: AY196923

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<400> SEQUENCE: 30
gctatggaaa ctatggcatt ctctcgattt tctatatgtt tttgtgtcct tctcccttgc 60
catggttctt tggctcagat atttagtcta ggcataaaatc catggcaaaa tcctcgacaa 120

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gggggttcta gggagtgttag gtttgatagg ctccaaagcgt ttgagccgct taggaaagtg	180
aggcatgaag ctggggttac agagacttt gatgagaaga atgagcgtt ccagtgcacc	240
ggtacattag taattcgctg cattatttag cctcagggcc ttctttacc tcgatactcc	300
aacactcctg gcctagata tatacccaa gggactgggt tactgggatt gacccctc	360
ggttgcggcag caacttacca aaagcaattt aggcatgggt gtcttgaagg aggaagccaa	420
aggcaaggaa aaaaattaag agatgaaaac caaaagatcc accaatttag gcaaggagat	480
gttggcac ttccttctgg tataaccacac tggttctata atgagggtga caccctgtt	540
gttgcttgtt tggttttga tgtaaacaac aatgctaattc aactcgaaacc aagacaaaaag	600
gagttcttgt tagctggtaa caatatacgag caacaagtgt ccaaccctc aatcaacaaa	660
cattctggcc aaaacatatt caatggattc aacactaagc tattaagtga ggccttaggc	720
gttaacatag aggtgaccag aaggctacaa agtcaaaatg accgaagagg agatatcatt	780
cgagtaaaga atggccttcg attgataaaa ccaactatca cacaacaaca ggaacaaaca	840
caagatcat accccaccaat tcaatatcat agagagcaac gatcaacaag caaatacaat	900
ggcttggatg agaacttctg tgcaatttagg gcaagggtta acatagaaaa ccctaattcat	960
gtctgatactt acaaccctcg tgctggagg attacaaatc tcaatagcca gaagttctcc	1020
attcttaacc ttgtccaaat gagtgctaca agagtaaattc tataccagaa tgctattctc	1080
tcaccattct ggaatattaa tgctcacagt ttgggtgtata caattcaagg gctgtctcg	1140
gttcagggtt ttagcaacca tggaaaggct gtatataatg gtgttctcg tccagggcaa	1200
ttactaatta taccacaaaa ttatgtggtt atgaagaaag cagagcttga aggatttcaa	1260
tttatcgcgt ttaagacaaa cccaaatgcc atggtaaacc acatcgcccc gaagaactca	1320
gttctccgtg caatgcctgt ggatgtgata gctaattcatc atcgcatctc aaggcaggaa	1380
gctcgttagct tgaagaataa taggggagaa gagattgggt ctttcactcc tagatatcaa	1440
caacaaaaaa tccaccaaga gtactcaat ccaaacgaaa gtgagactca agagggtatt	1500
taagccc	1507

<210> SEQ ID NO 31  
<211> LENGTH: 264  
<212> TYPE: DNA  
<213> ORGANISM: Oryza sativa  
<220> FEATURE:  
<221> NAME/KEY: misc\_feature  
<223> OTHER INFORMATION: LOCUS: OSU43530 264 bp mRNA linear PLN  
05-FEB-1997; DEFINITION: Oryza sativa metallothionein-like type 2  
(OsMT-2) mRNA, complete cds.; ACCESSION: U43530

<400> SEQUENCE: 31

tttggaaagaa agatgtcgtg ctgcggagga aactgcggct gcggatccgg ctgccagtgc	60
ggcagcggct gcggaggatg caagatgtac ccggagatgg ctgaggaggt gaccactacc	120
cagactgtca tcatgggtgt tgcaccttcc aagggtcatg ccgagggggtt ggaggccggc	180
gccgccccccg gagcaggagc agagaacggg tgcaagtgcg gcgacaactg cacctgcaac	240
ccctgcaact gcggcaagtg aagc	264

<210> SEQ ID NO 32  
<211> LENGTH: 1065  
<212> TYPE: DNA  
<213> ORGANISM: Oryza sativa  
<220> FEATURE:  
<221> NAME/KEY: misc\_feature  
<223> OTHER INFORMATION: LOCUS: AK064310 1065 bp mRNA linear PLN

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04-DEC-2008; DEFINITION: *Oryza sativa Japonica Group* cDNA  
clone:002-107-B12, full insert sequence.; ACCESSION: AK064310

&lt;400&gt; SEQUENCE: 32

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acactaccca gctcaactgaa gtagcaacgc agctcatcag tgtcaagatg gctactacta      60
agcatttggc tcttgcatac cttgtccctcc ttagcattgg tatgaccacc agtcaagaa     120
ccctccatgg ttatggcccc ggaggaggag gtggaggtgg tgggtgggt gaaggaggcg     180
gtgggtggta cggtgatcg ggctatggtt ctgggttcgg gtatggtgag ggaggtggta    240
gtggaggtgc tgccggtgcc gggatgggc gtggtgaggc cggcggcgcc ggtgggtgg      300
agggtgtgg ctccggctct ggctatgggt ctggcaagg ctctggctat ggtcaggtg     360
ttgggtggtc tggcgggtat ggtatggtg gggaggtgg tggtgccaa ggtgggtgg      420
ctggcggtca cgggcaagggt tctggctatg gttccggata tgggtcggt gctgggtgg      480
ctcatggtgg tggttatggc agtggtgccg gtggcggtgg cggtgaggc caaggtggag     540
gtccggctc tggctctggc tctggatatg gtctggctc tggcgaggc aacggacacc     600
actaagtcata ttccctctat cagctagcta caatatagcc tggttata agtgaaccgt     660
gatcagtgtat gagtctctct cgctgctta caaagagott gtggattgt atcgtatag     720
caactgtgtac tggcttgccg tttcatcaca taaagttggc aaagcttagt aaataaaacg     780
acctttgtat ctcagatcat acttctgtta ttacacggagc tggatgtcac taagattgt     840
atgcaagaac atagatttaa cccttgcct agctagttgt ttacatagta agcatgcaga     900
atggacttgt taactctcggt tggatcttgcg ttgtaaaactg gactgtttct cagtagggct     960
gttagcttagt atttcttagat tccaacatgt tggcttacac ggtgttaaaa aaatattgt     1020
cgagattgtat agtagactac ccagataaag ttcaattgg tcagc                         1065

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&lt;210&gt; SEQ ID NO 33

&lt;211&gt; LENGTH: 1139

&lt;212&gt; TYPE: DNA

<213> ORGANISM: *Oryza sativa*

&lt;220&gt; FEATURE:

&lt;221&gt; NAME/KEY: misc\_feature

<223> OTHER INFORMATION: LOCUS: AK064485 1139 bp mRNA linear PLN  
04-DEC-2008; DEFINITION: *Oryza sativa Japonica Group* cDNA  
clone:002-111-B07, full insert sequence.; ACCESSION: AK064485

&lt;400&gt; SEQUENCE: 33

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catcttact aaagaagatg gtaaaatccc agataagcgt gtcacagacc ttcagttgaa      60
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gaaactgaaa gatggtcgga tagtgaattt gaatgtgcag tcagatgacc ctcttgcac     180
acttcaagaa gctttcgaga aggtcaatcc tagattggcc ttcaatattt agctcaaatt     240
tgatgacaat ttgtatgttcc aggaggaaga gtcacttgtt atcctccagg ccatctgaa     300
ggttgttttt gagtatgcca aggtcggcc tattttttc tctagttcc agcctgacgc     360
tgcacaggtt atgcgaaaaat tgcaaagcac ataccctgtt tacttttgc ctaatggagg     420
cacagaaatc tacgctgacg tttaggaggaa ctcattggaa gaggccatca agctatgcct     480
cgccagcggc atgcaaggaa tcgtgtcgga ggcacggaga atttcaggc accccgctgc     540
tgtaccaaag atcaaagagg ctaacctctc cctactgact tatgaaacac tgaacaacgt     600
accggaggcg gtgtacatgc aacatctgtat ggggggtgaac ggtgtatcg tcgacctagt     660
gcaggagatc accgaggccg tctctgagct catcaccgtc cggagccctg acctgaacgc     720
cgataatttgc agcaatgggg cagcaaaaga cggcccaacgc ccacatttct cggcgtgtga     780

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aatctcattc ttgctgaggc tcatccctga gcttgtgcaa taatcgatcc agttgccttg	840
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gagggttaca agtttacaaa acttttaat aggtttagt actagagcta agccctggc	960
ctgtttttta atttgcctg gggacgtttt gattcatgat ggatcatgga tgatgcattc	1020
tgttagcaagg agttgaaaca gttgttgtt ctgttgtatc tgtaagactg taacgacgac	1080
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<210> SEQ ID NO 34  
<211> LENGTH: 957  
<212> TYPE: DNA  
<213> ORGANISM: Oryza sativa  
<220> FEATURE:  
<221> NAME/KEY: misc\_feature  
<223> OTHER INFORMATION: LOCUS: AK101309 957 bp mRNA linear PLN  
04-DEC-2008; DEFINITION: Oryza sativa Japonica Group cDNA  
clone:J033033N23, full insert sequence.; ACCESSION: AK101309

&lt;400&gt; SEQUENCE: 34

ggttcgatcc aagaagcaga gaggagaggg attcttgttt tcttgatcat ccatggcg	60
gctgccccgt ggtctcgaac acctggggcg gcggtaacgac ttctacgcgg cgtaccactc	120
caacccggcg aacgtgctcg tccacggcg gtgcgtgtgg cccatctcc tcaccgcgt	180
gctcccgctc cggtaacgcgc cgccgctgcc gctgctccga ttctactgcc cgctctggc	240
ccagtagctc cccgtgcagc tcggcttccc cgtaacgcgc ggcgtggcg cgtactacgc	300
gctcatggac cgccgcgcgg ggcggccgcgc cgccgctcc tcgcgtgcgc ggtgggcgc	360
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gcggccggccg ggcgtgggtt atggcccggt gcaggccgtg gtgcacggcgc cgctgtcgt	540
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ccaagtgttc gatataatgc gagtgccgtt gtcgtgtt tttcgggtt ctgcatacggt	660
tgttcgggta cgagccgacg ccggggttct acaagcgcgt ccaggcgagg gtggccgcga	720
tgcacaacgg gccgcggca ccggccggcg cgccggagaa gaaggaggag gaggagaagg	780
agaacgtgag caaggccgacg caggaggaga ggcggccgaa ggattcgttag gtgtttcga	840
gggagaggtt gacagacagg agaggccagc ctgcggatgt tatgaacttg taatagctt	900
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<210> SEQ ID NO 35  
<211> LENGTH: 630  
<212> TYPE: DNA  
<213> ORGANISM: Oryza sativa  
<220> FEATURE:  
<221> NAME/KEY: misc\_feature  
<223> OTHER INFORMATION: LOCUS: AK107983 630 bp mRNA linear PLN  
04-DEC-2008; DEFINITION: Oryza sativa Japonica Group cDNA  
clone:002-135-E08, full insert sequence.; ACCESSION: AK107983

&lt;400&gt; SEQUENCE: 35

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gtcgccctcc tcgcggatcac ggccgcgcgg caggccggcg gacggccccc cgccggcg	180
cccaagatgg ccccgatgcc cgatcccccgc ggcggatccc gggccaccgc ccccgccgc	240

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tccgtctccg	ccccgacccgg	cgccccccgt	gcctccctca	cgtacaccgc	caccggccagc	420
tccgttgcgg	tcgcggccgc	ggteggccgc	gccatcgtgt	tctagatgga	tggatggatg	480
atttgatcga	cgcgttggtt	atacggcgag	attctttttt	gtgattctgt	tctatttaca	540
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atggaatata	ttcttcatt	tattattacc				630

<210> SEQ\_ID NO 36  
<211> LENGTH: 1412  
<212> TYPE: DNA  
<213> ORGANISM: Oryza sativa  
<220> FEATURE:  
<221> NAME/KEY: misc\_feature  
<223> OTHER INFORMATION: LOCUS: AK099918 1412 bp mRNA linear PLN  
04-DEC-2008; DEFINITION: Oryza sativa Japonica Group cDNA  
clone:J013116J24, full insert sequence.; ACCESSION: AK099918

<400> SEQUENCE: 36

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cccgccggcc	cccgcaaccc	cacccgtctcc	gtcgccggcc	gcacccgggg	cgccatctgc	180
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<210> SEQ\_ID NO 37  
<211> LENGTH: 2771  
<212> TYPE: DNA

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<213> ORGANISM: Oryza sativa
<220> FEATURE:
<221> NAME/KEY: misc_feature
<223> OTHER INFORMATION: LOCUS: AK100306 2771 bp mRNA linear PLN
    04-DEC-2008; DEFINITION: Oryza sativa Japonica Group cDNA
    clone:J023078G01, full insert sequence.; ACCESSION: AK100306

<400> SEQUENCE: 37

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<210> SEQ ID NO 38
<211> LENGTH: 692
<212> TYPE: DNA
<213> ORGANISM: Oryza sativa
<220> FEATURE:
<221> NAME/KEY: misc_feature
<223> OTHER INFORMATION: LOCUS: X83434 692 bp mRNA linear PLN
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    b1.; ACCESSION: X83434
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gggggggctc ccgccccgtcg ggggcctgtc gcaggccgt caggagcctc aactccgccc 180  
ccaccaccac cgccgaccgc cgccaccgcct gcaactgcct caagaacgtg gccggcagca 240  
tcagcggccct caacgcccggc aatgccgcga gcatccctc caagtgcggc gtcagcatcc 300  
cctacaccat cagccctccat atcgactgtc ccagcgtgaa ctaatccgt cgtatcgctac 360  
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<210> SEQ ID NO 39
<211> LENGTH: 2882
<212> TYPE: DNA
<213> ORGANISM: Oryza sativa
<220> FEATURE:
<221> NAME/KEY: misc_feature
<223> OTHER INFORMATION: LOCUS: AK070414 2882 bp mRNA linear PLN
        04-DEC-2008; DEFINITION: Oryza sativa Japonica Group cDNA
        clone:T023053H09 full insert sequence ; ACCESSION: AK070414
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<210> SEQ\_ID NO 40  
<211> LENGTH: 984  
<212> TYPE: DNA  
<213> ORGANISM: Oryza sativa  
<220> FEATURE:  
<221> NAME/KEY: misc\_feature  
<223> OTHER INFORMATION: LOCUS: AK103220 984 bp mRNA linear PLN  
04-DEC-2008; DEFINITION: Oryza sativa Japonica Group cDNA  
clone:J033122018, full insert sequence.; ACCESSION: AK103220

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<210> SEQ\_ID NO 41  
<211> LENGTH: 2439  
<212> TYPE: DNA  
<213> ORGANISM: Oryza sativa  
<220> FEATURE:  
<221> NAME/KEY: misc\_feature  
<223> OTHER INFORMATION: LOCUS: AK121856 2439 bp mRNA linear PLN  
04-DEC-2008; DEFINITION: Oryza sativa Japonica Group cDNA  
clone:J033102G02, full insert sequence.; ACCESSION: AK121856

<400> SEQUENCE: 41

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ggtctttca aggaagatga tcctgagttc accacctcg acgcccgtac cgacgtcact	1800
atcaaaccctt catcgaaaga aaccgtttag attcctgtca ctgagaattc cacgatttgg	1860
tgggagctcc ggggtcttgg atgggagggtg agtacggag cagagttcac tcctgtatgcc	1920
gagggtggat acacagtcat cgtgcagaaa acgaggaagg tgcctgcaaa tgaggaacca	1980
atcatgatag gcagcttcaa ggtggcgag ccaggaaaga ttgtgtcaac gatcaacaac	2040
cctgcatcaa agaagaagaa gtcctctac agatccaagg tcaagagcac cagtgtgtcc	2100
gtttgaggtt gcagctgcct gatcaccaga ttccaccacaa tggcagctga actcattccc	2160
tgtatggaa gaaacctttt ggtttggtt cttaattta ttgggtttgc tggtttggtt	2220
cacattttgtt atttggtaa ttaaaaacca aagttagt gttttgtga tagttggaaag	2280
gagagggttg atatgatata atgacatcgat gatggttgt tgaggccaga ggacaaaaat	2340
tgtggaaagg ctgaagaata tctgtgttctt tgttatatctg tctgtacatt gcatctctgg	2400
attctcatgg acatgttaaa tttagaagta ctgtgtcattc	2439

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<211> LENGTH: 536  
<212> TYPE: DNA  
<213> ORGANISM: Oryza sativa  
<220> FEATURE:  
<221> NAME/KEY: misc\_feature  
<223> OTHER INFORMATION: LOCUS: AK062758 536 bp mRNA linear PLN  
04-DEC-2008; DEFINITION: Oryza sativa Japonica Group cDNA  
clone:001-106-G10, full insert sequence.; ACCESSION: AK062758

<400> SEQUENCE: 42

atccctgcaca cagccatttc acttgctttg ctccacgcct ccacacctaaatcgctct	60
cattagtaact caattaagtgc gatcgccgag cgattcagcg cggctctcg accgatctct	120
ccgtttcgctc agatcgatcg atcgatcgat cagggctgttgc tattgcgcgc tgtagtacgt	180
gagaggatcg gagtcggagc agcgagcgag atttgttgaag ccaaattaag tagatcgatc	240
tcatatcgat catccttaat taatttatca ttggagctgggt gctgggtata tcgctgcgc	300
tgcggcgct catataatc gtcgcctatcg tccctctgcgt catctgcgcgc tagcttagct	360
gatcccttcc tgcctccgcg atcattttgc atacatgttag tatgtcagta tgtagtgcgt	420
caaatgtgat catttgcgcg cccgttggttt ttgggcacgg tgattgacga cagaaagaat	480
tactccotttc atattttaat gtatgacacc attgactttt taaccaacgtt ttgacc	536

<210> SEQ ID NO 43  
<211> LENGTH: 2026  
<212> TYPE: DNA  
<213> ORGANISM: Oryza sativa  
<220> FEATURE:  
<221> NAME/KEY: misc\_feature  
<223> OTHER INFORMATION: LOCUS: AK103306 2026 bp mRNA linear PLN  
04-DEC-2008; DEFINITION: Oryza sativa Japonica Group cDNA  
clone:J033125E13, full insert sequence.; ACCESSION: AK103306

<400> SEQUENCE: 43

gcgagccgcg cgagcgac acacacagag gaggaggccg cgccgcgcgcgac gcgagctcg	60
cctctgcgtg cgccgcgcgcg agggtcgccgc cgccgcgcgcg gccgtcgatc ggccgcgcgt	120
cgtatggggag ctgctactcc gcctacgcct cctcgccaa gctgcgcggc cgcatcgca	180
agatctccct cgtcatcccc gacccgggtcc cgcacgcgcg ggcgcgcctcg cgcgcgaagg	240
acggcgtcga tggcacggc gacgacgtga ggggtgggtgg tgggtggctgt gacgatggcg	300
gtgacgtcgt cgccatcgcg acgacgcacgg cggacgaggct cgccgcgcgg tacgtctgg	360
ggaaggagct gggggcgccc gagttcgcccc tgacgcggcg gtgcacgcac gcggcgaccg	420
gggaggcgct ggccgtcaag acgatccgga agcacaggccg cctggcgccg ccgcgggtga	480
ccgcggcgaa ggccggggcc ggcacgggg aggacgtgaa gaggagggtg gccatcatgc	540
ggcgcatgtc gtcggcgctcg tctgcgcgcg gggcgccgcg cgccgtcgcc gccgcgcgtgg	600
tgcggctccg cgaggcatgc gaggacgcgcg cgcacggctc cgtccaccc tgcacggac	660
tctgcgaggc cggcgacgtt ttcgacccca tctgtggcgcc cggacactac tccgacgcgc	720
ccggccccaatctccgc accatcgctcg acgtcgtccca gctgtgccac tcgaacgggg	780
tgatacacccg cgatctgaag ccggagaact tccgttgcgc gaacaaatcg gaggactcg	840
cgctcaaggat catcgacttc ggcctctcg ttttcttcaa gccaggcgac cggttacgg	900
agggtggggg gagcgctac tacatggcgcc cggaggattt gcccggccgac tacggccgg	960
agggtggacgt gtggagcgcc ggcgtcattc tctacatcc tctctgcggc gtcctccat	1020
tctggggaga caacgacgcg aagatcgccg acggatccct ccgcggccatc atcgactca	1080
acaggggagcc attgcccagg gtctccgcca acgccaaggaa cctcgtcagg aggatgcttgc	1140

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atcctaacc	atccacccgc	ctgacggcca	aacaaggttc	tgagcatcca	tggctgaaga	1200
acgcggacac	ggcgccgaac	gtgtcgctgg	gacgcggcgt	gcggggcgagg	ctgcagcagt	1260
tctccgcat	gaacaagttc	aagaagaagg	ctctcgagg	ggtggcgccg	aacctggccgg	1320
gggaggaggt	ggacaagtac	gtgcagatgt	tccaccacat	ggacaaggac	aagaacgggc	1380
acctgtcgct	cgacgaactc	tttgaaggcc	tccacatcaa	cggccagccc	gtccccgagc	1440
ccgagatcg	gtgtctactc	gaagccgogg	acacggacgg	gaacgggacg	ctggactgog	1500
acgagttcg	gacgggtgtcg	gtgcacctga	agaagatgag	caacgacgag	tacctggcg	1560
cgccgttcaa	ctacttcgac	aaggacggca	gccccgttcat	cgagctggac	gagctgcggg	1620
aggaggtggg	cccaaaacgag	caggccatcc	tggagatct	ccgcgcacgtc	gacaccgaca	1680
aggacggccg	catcagctac	caggagttcg	agctcatgtat	gaagtcggc	gccgactgg	1740
ggaacgcctc	caggcacttc	tccagggcca	acttcagcac	cctcagcagg	aggctctgca	1800
aggatactct	tactccctga	tgatcgatga	tgcccaacaa	tctcatcaac	tttgtttcat	1860
caaggcttat	aaactgcaca	tctttcacct	tcaccagaca	gggataacta	gagagagata	1920
caagaccaac	tttagagaaat	gtattggagt	ggcaacacgca	tgcgatgtcg	cgcttggatt	1980
gtgaaatgg	aatggaaact	gaaatgggga	tcgtacccgt	tttgc		2026

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<210> SEQ ID NO 44
<211> LENGTH: 1154
<212> TYPE: DNA
<213> ORGANISM: Oryza sativa
<220> FEATURE:
<221> NAME/KEY: misc_feature
<223> OTHER INFORMATION: LOCUS: AK061207 1154 bp mRNA linear PLN
        04-DEC-2008; DEFINITION: Oryza sativa Japonica Group cDNA
        clone:006-210-C03. full insert sequence : ACCESSION: AK061207
```

<400> SEQUENCE: 44  
atatcgatct gatctctcta tatattttat ggattacctc tctgtttta agggaaagagc 60  
tagcgatcg a gtcggtag tatacgtca ctttgggtt ctcatgact ggcttaattt 120  
ctgaagatga gtaatctgt caaggctaa agctgggtac cagaggagaa agctgcagecg 180  
actgcttctg atgaacagaa tgacaagata aagaaagtcc gggactcct gggaaagccag 240  
atgacagcag agatgccgtc attttgtcg gacgccacca tccggccctt cctccggca 300  
aggaactgga gcacggagca agcaaccaag gctctaaag agactgtcaa atggaggcgt 360  
cagtcacggc cggacacaat ccgctggaa gacattctg gaaggagca cgaagctagg 420  
agaacatata tagccgacta cttgataag aatggaagga tcgtcttcat atcgaatccg 480  
acaattaaga gtaaaatcatc caccaaggac cagataaaac agtttagtgta taacctggag 540  
attttgcca tgcactcaga aaacatggaa gacgaatgtc ctgtttgggt aactgacttt 600  
caaggctggg tgctaacaat tacgccattt ccgttgcctc gtgaatgcac tcacataatt 660  
caaaaccatt atccagggtc gatttctgtc gcaatccca gcaacccacc aaggattttc 720  
gaatccccc ggaaggatgt gtgcattttc attgagccaa agttgaaaga aaaagtgaag 780  
ttcgttatata ctaacaatcc agagagccac aagatagttt ctgatatgtt tgattggac 840  
aagctggagt ctgcattttt gggggaggaac acactccat ttgacatggc caagtatgca 900  
gagagaatga aacgaagtga ccaaatgaga ggagctccca tgcacatggc tggctactct 960  
tgctctaccc aaacctgacc acataaaagt tcacccat tttttttttt ttcataaaaa 1020  
gtggcataaa tataatatacg ccggtagat ctgatataat gatgcatttc acttttaggt 1080

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tctattgtat ttatgttat actagaatta gggaaagatt aaggcgttag aagaaatcg 1920
aggaattccg gagttatcg tgatccttt ctatttctta tactttgtta tttgctttaa 1980
tagaaatatac atttcaagt 1999
```

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<210> SEQ ID NO 46
<211> LENGTH: 822
<212> TYPE: DNA
<213> ORGANISM: Oryza sativa
<220> FEATURE:
<221> NAME/KEY: misc_feature
<223> OTHER INFORMATION: LOCUS: NM_001064530 822 bp mRNA linear PLN
 08-JUN-2010; DEFINITION: Oryza sativa Japonica Group Os06g0598500
  (Os06g0598500) mRNA, complete cds.; ACCESSION: NM_001064530
```

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<400> SEQUENCE: 46
atgtcaccac aaacagaaac taaagcaagt gttggattta aagctggtgt taaggattat 60
aaattgactt actacacccc ggagtacgaa accaaggaca ctgatatctt ggcagcattc 120
cgagtaactc ctcagccggg gggtccgccc gaagaagcag gggctgcagt agctgccaa 180
tcttctactg gtacatgcga agaaatgatt aaaagagctg tatttgcgag ggaatttaggg 240
gttcctattt taatgcattt caacttaacc gggggattca ccgcaaatac tagttggct 300
cattattgcc gcgacaacgg cctacttctt cacattcacc gagcaatgca tgcagttatt 360
gatagacaga aaaatcatgg tatgcatttc cgtgtattag ctaaaggattt gcgtatgtct 420
ggggggagatc atatccacgc tggtagatgtt gtaggttaagt tagaagggga acgcgatatg 480
acttttagtt ttgttgattt attgcgcgtt gattttattt aaaaagatcg tgctcgccgt 540
atcttttca ctcaggactg ggtatccatg ccaggtgtta taccgggtgc ttcaggggt 600
attcatgttt ggcataatgcc agctctgacc gaaatctttt gagatgatcc tggatgtt 660
tttggtagatc gaaatctttt acatccttgg ggtatgcac ctgggtgcagc agctaattcg 720
gtggcttttag aagcctgtgt acaagctgtt aacaagggcg cgatcttgcgtt cgtgaaggta 780
atgaaattat ccgatcagct tgcaaatttggaa gtcctgaact ag 822
```

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<210> SEQ ID NO 47
<211> LENGTH: 547
<212> TYPE: DNA
<213> ORGANISM: Oryza sativa
<220> FEATURE:
<221> NAME/KEY: misc_feature
<223> OTHER INFORMATION: LOCUS: AK119900 547 bp mRNA linear PLN
  04-DEC-2008; DEFINITION: Oryza sativa Japonica Group cDNA
  clone:002-180-F02, full insert sequence.; ACCESSION: AK119900
```

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<400> SEQUENCE: 47
aaaaaaaaaaaa aaaaaaaaaaa acttggctgt tccgctgtccagcgctcg gctcgagcc 60
gccccgtcgaca ccgggtccga ccgttggcgt gacggcggc agccacctcg tcggctccgg 120
tggccacagc tacttggacg tcaggacaga agaggaattc aagaagggac atgtggagaa 180
ttctcttaat gtgcattcc tcttctttac ccctcaaggaa aaggaaaaga acacaaagtt 240
catagagcag gtggcattgc attatgataa ggaggacaac ataattgtgg gttgcataag 300
tggagtaaga tctgaactag catctgccga tctcatagca gccggattca aaaatgtaaa 360
gaacatggaa ggagggttaca tggcatgggt ggaaaatggc cttgcgggtga ataaacctct 420
atgtcaagaa gagctcttagt ttcatgttattt gtatTTTGTGATAATGACTT tcagttattt 480
tatgactggg tggtaatttgg atttgttagaa agggaaaggta taagaaaaca atgttatatt 540
```

## US 9,353,379 B2

89

90

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attcccc

547

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<210> SEQ_ID NO 48
<211> LENGTH: 1803
<212> TYPE: DNA
<213> ORGANISM: Oryza sativa
<220> FEATURE:
<221> NAME/KEY: misc_feature
<223> OTHER INFORMATION: LOCUS: RICRTUB1X 1803 bp mRNA linear PLN
  16-MAY-1995; DEFINITION: Rice beta-tubulin (RTUB-1) mRNA, complete
  cds.; ACCESSION: L19598

<400> SEQUENCE: 48

tcacccacc cctcgatctc tcgctcgccg ccggcgatcg gatcgctgg ttggatcatc      60
acaactcgcc aaagatgaga gagatcctgc acatccaagg cgggcaatgt ggcaaccaga     120
tcggtgccaa attctggag gtggtgtgtg atgagcatgg cattgaccct actggggcggt     180
acacaggcaa ctcaaatctg cagttggagc gtgtcaatgt gtactacaat gagggctccct   240
ggggtcgttt tggcttcgt gctgttctca tggaccttga gctggtaat atggacagtg     300
tcggcaactgg accctatggc cagatcttc ggcctgacaa cttgttctc gggcaatctg     360
gtgctggtaa caactgggcc aaggacact acactgggg tgccgagcta attgatctg     420
tttagatgt tggagggaaag gaggctgaga actgtgactg cttgtcaagg ttccaagtt     480
gcacatctct tgggtgttgt actggatctg gtatgggtac actgttgcata tcaaagatca   540
gggaggagta ccctgaccgc atgatgctga cattctcagt ttcccctca cccaaagtt     600
ctgatactgt ggttggccatacatgta cactctcagt ccatcgttg gttgagaatg     660
ctgatgagtg tatggtttgt gataatgagg ctctctacga cattgttctc aggactctca   720
agctgaccac acctagctt ggggatattga accattttgtt ttctggccacc atgagtggag  780
tcacatgctg octaagggttc cctggtcagt tgaactctga cctccgtaaat ctggcagtga  840
accttatccc ctteccctgt ctccacttctc tcatggtcgg attcggcccg ctgacatcac   900
gtggctccca gcagttccgt gcccttactg ttctgtgact cacacacgac atgtggatg  960
ccaagaacat gatgtgcgtc gctgttcctc gccatggccg ttacctcacc gcctctgcca 1020
tggccgtgg aaagatgagc accaaggagg ttgatgagca gatgatcaat gtccagaaca 1080
agaactcata ctacttcgtc gagttggatcc ccaacaatgt gaagttcagt gtctgtgaca 1140
ttccaccgag aggcccttccatc atggcatacga ctttcattgg caactcaaca tccatccagg 1200
agatgttccg gagggtgagc gaggcgttca ctgttatgtt caggagggaaat gctttctgc 1260
actggtagacac tggcgaaggc atggacgaga tggagttcac cgaggcagag agcaacatga 1320
acgacccctgt ctctgtgtac cagcgttacc aggtgtccac cggccgttgc gaggccgtt 1380
acgaggacgca ggagcgttccatc gaggcttgcg acatgttgcg tggcttttc ttgggtttc 1440
tagggccaggg ttttgggttgc ttgggttttc cgtcttacat tatcaccgtt ttaccgcctc 1500
gtacggccacc gccgggttccatc atgttttcgtc ttgggttttc cgtcttgcgtt atggggaccc 1560
ttttgggttgc tggatgttgc tggatgttgc gatatttcgtt gatgttgcgtt 1620
acctttccatc gttgggttccatc gttgttgcgtt ttgggttttc cgtcttgcgtt atggggaccc 1680
cgccgttccatc cgtggccatc tttgttgcgtt ttgggttttc cgtcttgcgtt 1740
gattgttgcgtt aatgttgcgtt ttgggttttc cgtcttgcgtt atggggaccc 1800
aaa                                         1803

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&lt;210&gt; SEQ\_ID NO 49

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<211> LENGTH: 568  
<212> TYPE: DNA  
<213> ORGANISM: Oryza sativa  
<220> FEATURE:  
<221> NAME/KEY: misc\_feature  
<223> OTHER INFORMATION: LOCUS: AK070851 568 bp mRNA linear PLN  
04-DEC-2008; DEFINITION: Oryza sativa Japonica Group cDNA  
clone:J023065B06, full insert sequence.; ACCESSION: AK070851

<400> SEQUENCE: 49

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gacacagctg cactacttgc actgaagtag ctagagcttg caagtaaaga tggctaccac      60
caagcatttt gtgttgctaa taatccttgt ctccttgc ataggaatga ccactagtgc      120
tagaactctc ttgggttatg gcattggagg agaagggtggc ggtgggtggag gaggagggtgg      180
tagcggcgga ggaggtggtg ggtatggtag cggctcaggc tatggatctg gcagcggtata      240
tggtaagggt ggccgggtgtt atgggtggagg atatggaagc ggtgggtgggt gtggaggcg      300
cggcggccaa ggcggaggat ctggctatgg ttctggctct ggctacgggt atggatccgg      360
aggagggtggt gggcactact agtcttatac tcggcaatgg agcaattcta ccgtggact      420
tgtgcactca tataatgtatga ttgtgttaaca ttggatataata cacaagcttg ttttcgtgtat      480
tatatgtataat gcttcttgat tgggtttgtat gctgccaat atatgcgtt cccaaatgtata      540
aataaaatcaa aaataaggtt gaaataacg      568

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<210> SEQ ID NO 50  
<211> LENGTH: 942  
<212> TYPE: DNA  
<213> ORGANISM: Oryza sativa  
<220> FEATURE:  
<221> NAME/KEY: misc\_feature  
<223> OTHER INFORMATION: LOCUS: NM\_001051294 942 bp mRNA linear PLN  
08-JUN-2010; DEFINITION: Oryza sativa Japonica Group Os01g0840300  
(Os01g0840300) mRNA, complete cds.; ACCESSION: NM\_001051294

<400> SEQUENCE: 50

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atggagacga cgacgacgac gttgggcggc ggcggcggcg ggcgggggg aggcttctcc      60
gatccggccgt ctccgcgttc gccggccgtg tcggccggct cggcggcgcc ggcggcgctg      120
gcgaacgcgc ggtggacgccc gaccaaggag cagatcgccg tgctggaggg gctgtacccgg      180
caggggctgc gcaacggcgc acggcggcgcg atacagcaga tcacggcgag gctccggag      240
cacggccaca ttgaggccaa gaacgtgttc tactggttcc agaaccacaa ggccggcag      300
cggcagaagc agaaggcagca gagttcgac tacttcagca agctgttccg ccggccggcc      360
ccgctgcggc tgctccacag gccactcgcg cggcccttcc ctctcgccat ggcgcggcagc      420
gegatgccac cgccggccgc gccggccggcg acgacgacga cggccgcgtg caacggcggt      480
ggtgtgtatgt tcaggacgccc aagttcatgt cgggtcgccgcaaaataacgc cagctactac      540
ccgcagcagc agacgcccgtt gctgtaccca gggatggaaag tggatggccca cgacaagtcc      600
acggcgcagc caccggccac caccacatg tacctgcagg caccgcggag cagcgcacac      660
ctcgccggcg cggctggccg cggcggcgccg gaagcggaaag gccatggccg ccgcggccgc      720
ggcgccgggtg ggccgcggac cctccagctg ttccccctgc agccccacccctt cgtgtccgc      780
gatcacaagc cgctccgcgc cgggagcgcgc tgccggccgc tgccccccgac gacgcgtcc      840
cgctccgcgt cggttctcggt ggagtcggag agctcggaca gccccagcag cgaggccct      900
cggttctacg acttcttcgg cgtccattct ggaggccgcgt ga      942

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<210> SEQ ID NO 51  
<211> LENGTH: 1152

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<212> TYPE: DNA  
 <213> ORGANISM: Oryza sativa  
 <220> FEATURE:  
 <221> NAME/KEY: misc\_feature  
 <223> OTHER INFORMATION: LOCUS: AK059805 1152 bp mRNA linear PLN  
 04-DEC-2008; DEFINITION: Oryza sativa Japonica Group cDNA  
 clone:006-204-G07, full insert sequence.; ACCESSION: AK059805  
  
 <400> SEQUENCE: 51

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cagctttccac tccacgaaaa ccctctgcct cttcgctcg ctgcaccctc gtctggcgta 60
tccatccatg gctgccagga agttcttctgt cgccggcaac tgaaaatgca atggacagg 120
ggaggacgtg aagaagatcg tcaccgtctt caacgaggcc gaggtgcctt ccgaggacgt 180
cgtcgagggtg gtggtaggcc cgccgttctgt ttccctgcgtt caggtgaagg gtttgcctcg 240
gccggacttc tccgtcgccc cgccagaattt ctgggtgcgc aaggccggcg ctttactgg 300
cgagatcagt gcccggatgt ttggtaacctt gcaggtgcctt tgggtgattt tgggacactc 360
tgagccggaga gcgctgtatgg gccaatcaag tgatttttttt gctgacaaaaa ttgcgtacgc 420
actttccaa ggtatcaagg taattgtttt cattggtag acccttgaac agagagaagc 480
aggaacaacg atggaaatgtt ttgcagcgc aactaaatcg attgcagaga agatatccga 540
ttggaccaat gttgttttgg catatgaacc agtttggcc attggacccg gcaagggttgc 600
aacccctgtt cagggtcagg aggttcatgtt tgggtctgaga aagtggctt tgactaatgt 660
tagtcctgca gttgtgttcaat caaccaggat tatattacaga ggctccgtaa atggagcaaa 720
ctgcaaaagaa cttgtgtctt aacctgtatgt tggatggattt cttgtttggag gagcttcatt 780
gaaggctgaa tttgtggaca tcatcaagtc tggatccgtc aagtcttctt ctttgttcc 840
tgggttgcac ccagatgtatgtt gtagggatttt atgctgcacaa ttttaatgtt tgacatgttt 900
gaccagcttgc ctttgttata tctcgttgc agtgtactcc atatcggtcc atagagcatg 960
cagccacccgtt tgggttgcctt ttttctttt gacttttttcccgagagga tcagatgttgc 1020
tgaaagtgttgc gttaatgttctt gtattatcg aagtttggatggatggatgtt gctataatag 1080
ttgttgcacccgtt tacttcgttgc gtgttgcactt gtcctttgttca tgggttgcattt 1140
gttttgcgttgc tc 1152

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<210> SEQ ID NO 52  
 <211> LENGTH: 1833  
 <212> TYPE: DNA  
 <213> ORGANISM: Oryza sativa  
 <220> FEATURE:  
 <221> NAME/KEY: misc\_feature  
 <223> OTHER INFORMATION: LOCUS: AK106244 1833 bp mRNA linear PLN  
 04-DEC-2008; DEFINITION: Oryza sativa Japonica Group cDNA  
 clone:002-100-D03, full insert sequence.; ACCESSION: AK106244

<400> SEQUENCE: 52

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ccacccctcccc caccggccatc gccattggccg ccgacggagga agaggagaag cttcgagtgg 60
agcgatctcg atggacccgtt gcccgttctgtt gcccggatgtt gtccggcaacc tctcgctgaa 120
gtatggccgtt gcccggccccc ccggccggagc cggggccgggg gtgcacccat ccacccgttcc 180
gtgttactgc aagatccggcc tcaacaatcg gcccgttccatcc accggccgacg cggccgtgtt 240
gtgttccggccccc tccggccggagg catcgccggcc gcccggccca gcccggccga cggggccgctt 300
ccggccggccgtt ttccacccctt ccaaggccgaa cctcgacccgc cttccacccgttcc accggccgtgtt 360
gttccgggtcg cgcacggccgaa ggctgttgc gtttgcgttgc gtttgcgttgc gtttgcgttgc 420
gttccgggtcg cgcacggccgaa ggctgttgc gtttgcgttgc gtttgcgttgc gtttgcgttgc 480

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<210> SEQ ID NO 53
<211> LENGTH: 954
<212> TYPE: DNA
<213> ORGANISM: Oryza sativa
<220> FEATURE:
<221> NAME/KEY: misc_feature
<223> OTHER INFORMATION: LOCUS: AK108127 954 bp mRNA linear PLN
        04-DEC-2008; DEFINITION: Oryza sativa Japonica Group cDNA
        clone:002-139-D06, full insert sequence.; ACCESSION: AK108127
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<400> SEQUENCE: 53

acaaccagca gcgcgagttac accccaaatcca gcagcatcgta cgattcggccg agctaggggc 60  
agaggcagaa gtgcagaacg tacacgcgtg gtgagcagag gagcaatggc gactcggctt 120  
ctgtctgtgc tgctgtctt gctcggcatg tcgctgaaaatgatcagaggg ggcgtgggtc 180  
gtgtgcaggc cggacgtggc ggaggcgccg ctgcagaagg cgctggacta cgcgtgcggg 240  
cacggcgcgg actgcgcgccccc ggtgacgcggc agcgggtcggt gctacagcccc aaacaacgtg 300  
ggggcgcact gtcctacgc cgccaaacagc tacttccagc ggaattccca ggccaaggggc 360  
gccacactgcg acttcggccgg cgccgcaccacc ctctccctcca ccgaccccaagc 420  
tgcaaataacc ctgcaaccgc aagtgtcgca gggacaagcgcc acggaaacccgg cacggcgggt 480  
gcaggcacag gcaccggtagc aagcacgagc acgagcacgaa gcacttcttc cccgggtct 540  
tcaactdqcaq ccacqqqtac qccqatcatq qqadqqqacqt tcqctacqcc qatcqccqqc 600

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ggcgcgctcg	ggccgacgac	tagegccttg	aatcctgaca	gcagcgaagc	accgtccccg	660
tcccttggac	gtcatttct	cctcacgtgc	attgcctcca	tgctgtctc	taattttctt	720
ttggcgtagt	aatttggccg	gaaagtaacg	tgaactgaag	cagcatactg	tgggagaatg	780
gagatttggc	tttcctttc	tgcttggag	cttgtcggg	cctgttctt	ttctttacca	840
gaactgcagg	tcctgggtgt	ttacacctgt	tgtaccatgt	cccagtgaac	tggccgtttg	900
catcatggca	tgttagtat	gttacgtgc	tagtgcagc	acatttcac	tgtat	954

<210> SEQ ID NO 54  
<211> LENGTH: 1100  
<212> TYPE: DNA  
<213> ORGANISM: Oryza sativa  
<220> FEATURE:  
<221> NAME/KEY: misc\_feature  
<223> OTHER INFORMATION: LOCUS: AK061237 1100 bp mRNA linear PLN  
04-DEC-2008; DEFINITION: Oryza sativa Japonica Group cDNA  
clone:006-211-C03, full insert sequence.; ACCESSION: AK061237

<400> SEQUENCE: 54

atcagatcaa	agcaagcaaa	caacaacaaa	aaccacttct	cgtgggtaga	agagagagag	60
agcgaggcga	tttcgaccaa	gaagatggcc	gggatcgtgg	tgggtttcga	cttcgacaaag	120
acgatcatcg	acgtcgacag	cgacaactgg	gtcggtggacg	ggctcggcac	gacggaggag	180
ttcgagcggc	tgctgcccac	catgccgtgg	aacaccctca	tggacaccat	gatggggcag	240
ctccacgcga	gcccgaagtc	gtcgccgac	gtcgccggcg	tgctcaggtc	ggcgcgcgctc	300
gaccgcgcgc	tcgtgcgcgc	catcaaggcc	tgctacggcc	tgggctgcga	cctccggatc	360
ctcagegacg	ccaaccgctt	cttcatcgac	accatctcg	accaccacgg	cctcacgggt	420
tacttctccg	agatcaacac	caaccggcgc	gccgtcgacg	ccgcccacccg	ccgcctccgc	480
atcgccgcgt	accacgactt	ccacgcggc	ccgcacgggt	gcccgttcgg	gatctgccc	540
cccaacatgt	gcaaggccca	ggtgcgtgcac	cgcacccgc	cctccgcgg	cgccgcggc	600
aagagggtca	tctacccctgg	tgacggccgc	ggggactact	gcccgtcgct	ccgcctccgc	660
cgcgacgact	tcatgtgcc	acgcggggc	ttcccccgtgt	gggagetcat	ctgcgaggac	720
ccgtcgctgc	tccacgcgg	ggtgcactcg	tgggcccacg	gcccgcgat	ggaggagacg	780
ctgctcgccgc	tggteggcag	ggtgcgtcctc	gaggagagg	acctgcgcgc	gctcgactgc	840
aagctcgagt	cgttgccggc	cgtcgccgt	caggacggca	tgcccatgac	gctccggatc	900
aagaactgtat	aatggccgc	gacgaacgt	cgcacgtt	cgagcgcgaa	acggctagct	960
cgaacaatgt	gtgtgtgagg	attgcgatac	gggtataatt	ttaactatgt	actgatttt	1020
cgctacgcgc	tgattgagcc	tgcgattagt	agaggctcat	tgtatcttgc	ccgatcaatt	1080
gaagtaaaac	atttggcttg					1100

<210> SEQ ID NO 55  
<211> LENGTH: 459  
<212> TYPE: DNA  
<213> ORGANISM: Oryza sativa  
<220> FEATURE:  
<221> NAME/KEY: misc\_feature  
<223> OTHER INFORMATION: LOCUS: AK062517 459 bp mRNA linear PLN  
04-DEC-2008; DEFINITION: Oryza sativa Japonica Group cDNA  
clone:001-104-C09, full insert sequence.; ACCESSION: AK062517

<400> SEQUENCE: 55

ataagcaagg	gaaatatcag	catctgcaaa	aatcatctca	gattcggtcg	ccatggcgag	60
gatcccggtc	gccgcgcac	tcgtgcgcac	cctcttttc	gcacatgcgcg	cggtgcac	120

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ggcgccccggc cggtctccca ccagcgacgg gacgtcggtg gatcaaggga tcgcataacct 180  
 gtgtatgtc gtggcgctgg tgctcaccta cctcatccac cctctcgacg cctccctccgc 240  
 ctacaagctc ttctgagctt atgcagagat ctcttcgtcg ccatgggttt ccttctcctt 300  
 ctggatctc ctccctccctcc cttttgatag tctagtgggtg gatctctcat tctcggtgta 360  
 attaattatgt gggattttta tattttttt cagctcgctg tcgtttgtaa tttgggtcg 420  
 tggtaactcgctg ctgaggatcg atttcgattt gtgtatatac 459

<210> SEQ ID NO 56  
 <211> LENGTH: 3707  
 <212> TYPE: DNA  
 <213> ORGANISM: Oryza sativa  
 <220> FEATURE:  
 <221> NAME/KEY: misc\_feature  
 <223> OTHER INFORMATION: LOCUS: AK065259 3707 bp mRNA linear PLN  
 04-DEC-2008; DEFINITION: Oryza sativa Japonica Group cDNA  
 clone:J013002J18, full insert sequence.; ACCESSION: AK065259

<400> SEQUENCE: 56

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 ctccctctcc tcttcttctc caccacctcc attgctgctc gcctctctca cctccctctc 180  
 ctccctttgt gtggagctcg tcggcgctga ggttagctta gctagctaaat cttgtttgc 240  
 cattgttgttgc ttcttggtgt tcggagaggg agcttgctt tgcccttgagg ggagaggcaa 300  
 aggcatcgc aatggcgcca gcggtggccg gcccggagg gaggaggaaat aatgggggg 360  
 tgaacgggaa cgccggggcg ccggcggtcg tgtgcgggtt cccgtgtgc gcgtgcgcgg 420  
 gggcgccggc ggtggcgctcg cgccgcgtcg cggccgcacat ggacatcgat gcccgggggc 480  
 agatcgccgc cgtcaacgcgac gagagctggg tcgcccgtcgat cctcagcgac agcgacgc 540  
 ccccccggcgc cggcgacgtc caggcgccccc tcgacgaccg ccccgcttcc cgtaccgaga 600  
 agatcaaggcg cgtccctctc cacccttacc gggtgctgat ctttgtgagg ctgatcgatgt 660  
 tcacactgtt cgtatgtatgg cgtatcgacg acaagaaccc ggacgcgtatg tggctgtggg 720  
 tgacgtcgat cggccggcgag ttcttggtcg ggttctcgat gctgctcgac cagctccca 780  
 agctgaaccc gatcaaccgc gtcggcgacc tcgcccgtctt ccggccgcgc ttcgaccacg 840  
 ccgacgggac ctccctccctc ccggggctgg acatcttcgtt caccacccgc gaccgatca 900  
 aggagccat cctgtcgacg cgcaactcca tcctctccat cctcgcgcgc gactaccccg 960  
 tcgaccgcaa cacctgtac tcctccgtcg actctggat gtcctcacc tacgaggcca 1020  
 tggcgccggc ggccaaatggc gcgacgtgtt gggtgccctt ctgcggaaag caccgcgtcg 1080  
 agccgcggcg gcttgagagc tacttcgacg tcaagtccca cccctacatg gggagggcgc 1140  
 aggaggagtt cgtcaacgcgac cggcgccgcg tccgcaaggat gtacgacgac ttcaaggcca 1200  
 ggatcaacgg ctcgagcac gacatcaacg agaggtccga tcctacaac gcccgcggcg 1260  
 ggcgtcaagga cggcgagccc cggccgcaccc ggtggcccg cgggtcgac gggaggcg 1320  
 cctggatcgatcgac gcaatcgatcgac aaccaccgca agggcgacca cggccggatc gtcctgggtgt 1380  
 tgctgaacca cccgagccac gcacggcagc tggggccgc ggcgagcgcc gacaaccgc 1440  
 tggacttcgatcgac gggcgatggac gtggcgctgc cgtatcgatgtt gtacgtcgatcgatcg 1500  
 gccccgggtg caaccaccag aagaaggccg gcccgtgaa cggcgctgacc cggccctccg 1560  
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cccaggcgct ccgcgcggc atctgcttca tgctcgccg cgacagcgac accgtcgctg	1680
tctgtccattt cccgcagcgc ttgcaggcg tcgacccac cgacctat gctaaccaca	1740
accgttatctt ttgcacggc acgtccgtg ccctcgacgg gtcgcagggg ctatctacg	1800
tccggcaccgg gtgtctcttc cgccgcatac cgctgtacgg ttgcagccg ccgaggatca	1860
acgtcgccgg accgtgtttc ccgaggctcg gtggatgtt cgccaagaac aggtaccaga	1920
agcctgggtt cgagatgacc aagctggtg ccaagccgtt ggccgcggc ccggggcgca	1980
cggtgtggaa ggggaagcac gggttctgc cgatgccaa gaaggctac ggcaagtctgg	2040
acgcgttcgc cgacaccatc ccgcgcgcgt cgacccgtc ggcgtacgg gcggaggcgg	2100
cggtgtggggc cgacgaggcg gcatcgccg aggccgtat ggtacggcg gcccgtacg	2160
agaagaagac cgggtggggg agcgcacatcg ggtgggtgtt cggcacggtg acggaggacg	2220
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cgcacgcgtt catcgggacg ggcgcgcatac acctgacggg gaggctgttc caggtgtcc	2340
ggtgttgcac gggttcgctg gagatcttct tctcgaggaa caaccgcgtt ttggggagca	2400
cgttcctgcg cccgcgtcag cgctggcgat acatcaacat caccacctac ccgttacgg	2460
cgtgttctt catcttctac accaccgtgc cggcgctgtc gttcgtacg gggcacttca	2520
tctgtcagag gccgaccacc atgttctacg tctacctgcg catcggtc gggacgtgc	2580
tcatctcgc cgtgtggag gtgaagtggg cgggggtcac cgtgttgcag tggttcagga	2640
acgggcgtt ctggatgacg gccagctgtt cgcctaccc cgccgcgtt ctgcagggtt	2700
tcaccaaggt ggtgttccgg cgggacatct cgttcaagct caccttcaag ctccccccg	2760
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tgcgtcgcgg cgagtgacg cactggctca aggtcgccg cggcggttcc ttcaacttct	2940
gggtccctt ccacctctac cccttcgcgca agggcatctt cgggaagcac ggcaagacgc	3000
cgggtgtgtt gtcgtctgg tgggccttca cttcgatcat caccgtgtt ctctacatca	3060
acatccccca catccatggc cccggccgc acggcgccgc ctcaccatcc cacggccacc	3120
acagcgccca tggcaccaag aagtacgact tcaacctacgc ctggccatga ggacgcgtt	3180
gccggagacg aagaagaaa cacaacaag aacaagacg cacaacaac accaacaaca	3240
acaaacacga gatgatcgtt ttctactaca cgtgtgtcaca acaacacata ctactgaaca	3300
ctgtgcgtgc atttgatcga ggcacccgcg aaaatttggaa agttttttt cttttttct	3360
tttaaccttt tttttcttc ttttgcggc ctccctcttc tcttttttt tctttttgt	3420
tttgtccaga aaaaagatgg tttttttttt atttagttt ttaattacat gtggtaatta	3480
attatgtatt atacattact gcaaggaaga gaggggggtt tacaggtggg gcccgggggg	3540
tgggggtgtgg tttttttttt tactgtacat gctggggatgt tttttttttt tttttttttt	3600
agacaagagt cacagagagt gagagaaaga gaggctggaa gtggccccc ccaggtgggt	3660
gtgggtattct tttagtacat ggaaacaata aatthaattt cattatt	3707

<210> SEQ ID NO 57  
<211> LENGTH: 1323  
<212> TYPE: DNA  
<213> ORGANISM: Oryza sativa  
<220> FEATURE:  
<221> NAME/KEY: misc\_feature  
<223> OTHER INFORMATION: LOCUS: AK065604 1323 bp mRNA linear PLN

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04-DEC-2008; DEFINITION: *Oryza sativa Japonica Group* cDNA  
clone:J013034F24, full insert sequence.; ACCESSION: AK065604

<400> SEQUENCE: 57

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cgcgccgagc ggcggggagg agagattgga tcgacgcccgg ttagtttagga agggagcttg    180
gagagatggc ggcttcccgcg aggccccgtcg gctcgccggg ggagagggcg acgagctcg    240
ccatggcgtt cagcctgttc agccgctacg tccgccagaa cggccgcggcc gccgcggagc    300
tcggcctcggt catcagaggt gagggtgagg ctccgaggggc ggccgcggcg acgatgagct   360
tgctgccccgg ggaggccggag aggaagaagg agaccatgga gcttccccg cagacgcggc    420
gettggcca gcaggatgcc atcaccgcgg attctgtgc tgatgttagg gaacaagagc     480
ctgagaggcg tcaagtgacc atcttctatg gtggaaaggt gctcgtgtc aacgacttcc   540
cagccgacaa ggcaaaggcg ttgatgcggc tggctagcaa gggcagcccg gtggctctc    600
agaatgcgc ggcacactgca ccagcagctg ttacagacaa caccaaggcc cctatggccg   660
tgccggcccccc ggtcagtagc ttgctctacag ctcaggccggc tgctcagaag cctgtcgcc  720
cgaatgttttc tgatatgcct attgttagga aggcatactt ccacagggttc ttgagaaga   780
gaaaggatcg tcttaatgca aagacgccc accagggttc tcttcagat gcaaccccg     840
tcaagaagga gcctgagagc caaccatggc tggacttagg gcccgaacggc gtcgtgaagc  900
ccatagaacg cggccaatga ggttgcattgg aaacttcacc aaagctcttc aaaaatatat  960
gaaatgcgt tgccgttaga gtaccaaattt atatatgttc tctcataccc catgttttag 1020
tgttttttct ttagttcggtt cttctatgtt gttcactgtt cacatagttc tgaatgtaaa 1080
ggaaaaagggt agtagcattt acgaccatgtt ccgagagccct gagagaaggtt ttggcatgtt 1140
aacacatgtt gactctgaaa atggccatgc taagttttttt atttatttgc aaggaaacat 1200
atatgttatct tctgaaattt tgatcatgtt gctttgtcct tggctctacg cttgtctat 1260
gcaaaagcaaa gtgattgtaa tggattcaga atataattca tatttcattt gttgatttct 1320
gtt                                         1323

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<210> SEQ ID NO 58

<211> LENGTH: 1077

<212> TYPE: DNA

<213> ORGANISM: *Oryza sativa*

<220> FEATURE:

<221> NAME/KEY: misc\_feature

<223> OTHER INFORMATION: LOCUS: AK106964 1077 bp mRNA linear PLN  
04-DEC-2008; DEFINITION: *Oryza sativa Japonica Group* cDNA  
clone:002-119-F09, full insert sequence.; ACCESSION: AK106964

<400> SEQUENCE: 58

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actctcctcc attccttcac tgccttcact tccctgtca gtcagcagc tcagggactt      60
cgaggatgtt ggaatccaaa gcagccatga tggtgaccat cctgctctgc tgctctcca    120
tctcacccggc gttcgccagc aagcacaagg gtccgccccg cgcggccggcc gtaaggctcc  180
cgccgtcgcc ggcgcggcgtcc ccggccggcgcc cgcgcacgtt ggacctcgcc gacctcctga 240
gctggccggg tccgttccac acgttccctcg acctcctggaa gaagacggac gtgctcaggaa 300
cgttccagat ccaggcgaac ggcagcaagg acgggatcac ggtgttgc cccaggacg      360
ggcggttcgc gtcgtggcg aggtcgccgaa cggcgaacctt cacctccgac cagctcaagt 420
cgctggcgct gtaccacgcg cttccggaggt actactccctt cgcccgagttc aacaggctgg 480

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gccccggc cagcccggtg cccacgctcg	540
ccggcggcga gtacacggc aacgtcaccg	
acgacatggg caccgtccat gtcgggtcga	600
tgtggtccaa ccccaagatc agcagcagcg	
tctactccac ccgcggccgtc	660
ccgcgtctacg aggtggacag ggtgctctc	
ccgatgcaga	
tcttcaggac cgacccgccc atggcccggt	720
cgccggccgc ggcgcggac gccaaggccg	
cctccgacgc cgccagcccg ctccccggga	780
agtgcgtcag cgcccaaggcg aaggccgacg	
agaagaagag ctgcgtcgtc cgcgcgtcg	840
cgccggcatt gccggctact	
tcttggctct tgctgcatact gcctcagtg	900
gattgctgtc cctgtgttga tgctaagaaa	
cttttctaat tctttttttt tttgcatgga	960
tttgtggttt ctggataatt ttattcttg	
gacatagtag gggcattgtg ttaggattaa	1020
gttttatggg agtatttccg tcataatgctt	
gtatgattag tcgtatgacta cttgtttatc	1077
cgatgcgtgt tgcgaaattgc acaaggt	

<210> SEQ ID NO 59  
<211> LENGTH: 1347  
<212> TYPE: DNA  
<213> ORGANISM: Oryza sativa  
<220> FEATURE:  
<221> NAME/KEY: misc\_feature  
<223> OTHER INFORMATION: LOCUS: AK060983 1347 bp mRNA linear PLN  
04-DEC-2008; DEFINITION: Oryza sativa Japonica Group cDNA  
clone:006-203-A12, full insert sequence.; ACCESSION: AK060983

<400> SEQUENCE: 59	
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tccactgccaa	
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agaagacag	
gaagatgcac gccaagacgg actcggaggt gacgagcctg	180
gcccgcgcgt cgccatcgt	
gtccccggacg tcgcggcg	240
ggccggcggt gtactacgtc	
cagagccgt cgagggactc	
cgaggactc	300
gcacgtcggt	
ccggcgtcga	
gccccatggg	
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gcccacccaa	
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aggggtggca	
ggagatggc	480
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tcttcctctt	
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ccacagatag	
tcatcaagag	
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ggggacggac	
gcgtcgctgg	
tgccgacgga	
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cggtaagact	
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aacacggggca	
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cagctgacgc	
tggcgccgg	
cgatctcaac	840
aagtctacc	
aggccggag	
cagccggagg	
acggtgagcg	
tgggggtgat	
ggggaaacaag	900
gtccggctgt	
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cagcatggct	960
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tctgtgcatt	
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tttgccgatc	
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atttcccagt	
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attataccaa	
acgatcaatt	1320
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gtggcagaaa	
atgaaataca	
cagagctgag	1347
tatgtttatc	
tttcgag	

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<210> SEQ\_ID NO 60  
<211> LENGTH: 2714  
<212> TYPE: DNA  
<213> ORGANISM: Oryza sativa  
<220> FEATURE:  
<221> NAME/KEY: misc\_feature  
<223> OTHER INFORMATION: LOCUS: AK099719 2714 bp mRNA linear PLN  
04-DEC-2008; DEFINITION: Oryza sativa Japonica Group cDNA  
clone:J013088B06, full insert sequence.; ACCESSION: AK099719

<400> SEQUENCE: 60

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ccccagctac cacccctctc cctctctctc ctccatcagtc agctttgcct agatcacagc	180
tggatttctt ggccgcatec tcctcttctt ccaggtctct tcctatcgtc tagttgttc	240
ttcccttccc ttttttttgc ctttgtgcgc gtgaggatca gtgagctgtg cggctcttt	300
ttcttggtct tgctggactg ctgctgcctt aaaagatctt gcctttctt ctcactttcc	360
ccggaggaga gagagaggag atccacatct ctgatggttc ttggatctc ctaggagttt	420
tttctacctg tgctgctgtc gctgatttctt ttttcgttattt gctagcggtt gtgtggtcgg	480
agctcgctt cttgggtcga gatctcgaca cttcttgaga tggattttt caccgagttac	540
ggtgaggaa acaggtacaa gatagaagag gttataggaa aaggaggatc ttgtgtgttt	600
tgctctgctt tggacactca caccgggtat aaagttgtca tcaagaaaat caatgacatc	660
tttgagcatg tgcgtcgatgc aacacggata cttcgcgaga tcaagttgtc tagactcctg	720
cgcacatccgg atatcgatggaa aattaagcat attctacttc ctccatcaag gagagaattc	780
aaggatatctt acgtttttt tgaactcatg gagtctgtt tgcaccaagt tataaaggcc	840
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aaatatataac acacagttt ttgcacaaa ttttttttttcat cgagatctca aaccaaaaaa	960
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ggttcatcaa cttgtatataa ttacagatct actaggaaca ctttctccag aaacaatatc	1260
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gttagctttt gatccaaaag atcggccaag tgctgaggag ggcgttgcgtc atccttattt	1440
caagaacata gccaatgttgc atagagagcc ttctgtctca cctatcacaag agcttagat	1500
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aatccctcaa ggtgtgtctg caagacctgg taaagcgtt ggttcgggtgc tgcgttacgg	1920
taactgttgc acatccgttgc ctgagcaaca atatgacgtc cgaagggttgc ttccggaaacc	1980

agcaattgct ccaaacagca gtgttcctct gggagctca taccaggaa gaaaccagac	2040
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gccccatgtg gcgaacaaac tgccctgctac cgatcgatggt cggagtggcc actggtaat	2160
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cttggggct tttccatgtat aaaaacagta gaattgtact gcaagecccta gtggattagg	2340
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atgttggaca ttgcTTtagt atttcgttct caggaacatt gttcccttg cagagctagg	2640
agctgcaact atgtactact atctgacatt gctgtaaactt gtAAAactt attgcatttc	2700
aagtattttc octc	2714

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The invention claimed is:

1. A method for hydroponic cultivation of a genetically-modified plant comprising: cultivating the genetically-modified plant in a hydroponic medium,  
wherein the genetically-modified plant is transformed by  
introducing an expression vector comprising:  
a promoter that regulates expression of RNA expressed in  
a seed that satisfies the formula (1):

$$V/W > 1.0$$

(1), 35

wherein V and W are determined as follows:

V is an amount of RNA in the seed of a predetermined plant when the predetermined plant is cultivated in a hydroponic medium adjusted so that nitrate nitrogen is 70 mg/L to 750 mg/L and/or ammonium nitrogen is 70 mg/L to 750 mg/L for a period which starts from 30 days before the expected flowering date and ends at the flowering day, and W is an amount of RNA contained in a seed of the predetermined plant when the plant is cultivated in a medium adjusted so that nitrogen is 0 mg/L to 50 mg/L for a period which starts from 30 days before the expected flowering date and ends at the flowering day, wherein the predetermined plant is the same species as the genetically-modified plant, wherein RNA is extracted from the seeds 15 to 25 days after the flowering, 50  
wherein the promoter is a glutelin promoter; and a polynucleotide located downstream of the promoter and encoding a protein of interest, 55  
wherein the medium for cultivating the genetically-modified plant is adjusted so that a content of nitrate nitrogen is from 70 mg/L to 750 mg/L and a content of ammonium nitrogen is from 50 mg/L to 750 mg/L for a period

which starts from 30 days before the expected flowering date and ends at the flowering day of the genetically-modified plant, and  
wherein the ratio of the contents of the nitrate nitrogen to the ammonium nitrogen is from 3:1 to 1:3.

2. The method according to claim 1, wherein the predetermined plant is a poaceous plant and the genetically-modified plant is a poaceous plant.

3. A method for production of a seed comprising cultivating the genetically-modified plant according to the method of claim 1 and collecting the seed.

4. The method according to claim 3, wherein the plant is a rice plant and the seed is a rice seed.

5. A method for hydroponic cultivation of a genetically-modified plant comprising:  
cultivating the genetically-modified plant in a hydroponic medium,  
wherein the genetically-modified plant is transformed by  
introducing an expression vector comprising:  
a promoter that regulates expression of RNA expressed in  
a seed, wherein the promoter is a glutelin promoter; and  
a polynucleotide located downstream of the promoter and  
encoding a protein of interest,

wherein the medium for cultivating the genetically-modified plant is adjusted so that a content of nitrate nitrogen is from 70 mg/L to 750 mg/L and a content of ammonium nitrogen is from 50 mg/L to 750 mg/L for a period which starts from 30 days before the expected flowering date and ends at the flowering day of the genetically-modified plant, and  
wherein the ratio of the contents of the nitrate nitrogen to the ammonium nitrogen is from 3:1 to 1:3.

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